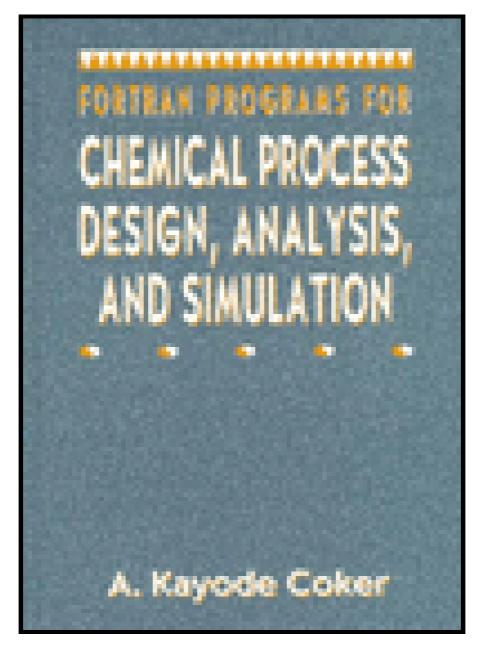
FORTRAN PROGRAMS FOR CHEMICAL PROCESS DESIGN, ANALYSIS, AND SIMULATION

A. Kayode Coker

Fortran Programs for Chemical Process Design, Analysis, and Simulation

by <u>A. Kayode Coker</u>



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A. Kayode Coker

Preface

The increased use of personal computers by engineers has greatly revolutionized and expedited the design of process plants and equipment in the chemical process industries. In addition, the availability of expensive flowsheeting packages has reduced to some extent the requirement of programming languages. However, situations often occur where either a package is not readily available, too expensive, or of a limited scope, and the design problem lies beyond the boundary of the flowsheeting packages. The designer is often required to review other alternatives or develop his or her own specific program for the application. This requirement is even more necessary when personal computers are used. A textbook analyzing the fundamental theory with equations, tables, and figures is not readily available to the designer.

My aim in writing this text is to present fundamental theory, along with computer programs, to solve a wide range of design problems. This text will provide consultants, designers, technologists, and practicing process engineers with structured programs that can be used to solve a wide range of process engineering problems and thus analyze and optimize process equipment on a regular basis. This book will also serve as a guide to practicing engineers and chemical engineering students who are interested in applying mathematical modeling and computer techniques to solve problems in process plant operations. This book is unique in that adequate information (theory, equations, figures, and tables) is provided to enable the reader to perform either hand calculations or to write alternative programs.

FORTRAN is still recognized as the most versatile programming language, even forty years after its introduction. It is a powerful general purpose language specially suited to complex, scientific, mathematical, engineering, and financial algorithms. The earlier versions of the FORTRAN language (FORTRAN IV, F66) could only be used on large mainframe systems that accepted programs in batch. Turnaround was slow; interaction was nil. The availability of microcomputers (IBM personal computers or compatible) has made it easier for numerical techniques, simulation, and design problems to be developed. Computer software can now be used interactively on such systems.

This book is unique in that to date no textbook on chemical process deign and simulation using microcomputers and the FORTRAN programming language has been published. FORTRAN 90 is a major development of the FORTRAN language and includes all the computer programs in this text as a subset. To assist the user and to demonstrate the validity of the methods, worked examples of practical industrial relevance are provided throughout the text.

A listing of all the programs is presented in the book, and source codes for these programs are provided on a floppy diskette. A FORTRAN compiler is required to compile and run these source codes. A menu driven software program for calculating the pressure drop of incompressible single-phase fluids (PIPECAL) is provided in Chapter 3.

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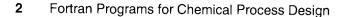
CHAPTER 1

Numerical Computation

INTRODUCTION

Engineers, technologists, and scientists have employed numerical methods of analysis to solve a wide range of steady and transient problems. The fundamentals are essential in the basic operations of curve fitting, approximation, interpolation, numerical solutions of simultaneous linear and nonlinear equations, numerical differentiation and integration. These requirements are greater when new processes are designed. Engineers also need theoretical information and data from published works to construct mathematical models that simulate new processes. Developing mathematical models with personal computers sometimes involves experimental programs to obtain the required information for the models. Developing an experimental program is strongly dependent on the knowledge of the process with theory, where the whole modification can be produced by some form of mathematical models or regression analyses. Figure 1-1 shows the relationship between mathematical modeling and regression analysis.

Texts [1-5] with computer programs and sometimes with supplied software are now available for scientists and engineers. They must fit a function or functions to measure data that fluctuate, which result from random error of measurement. If the number of data points equals the order of the polynomial plus one, we can exactly fit a polynomial to the data points. Fitting a function to a set of data requires more data than the order of a polynomial. The accuracy of the fitted curve depends on the large number of experimental data. In this chapter, we will develop



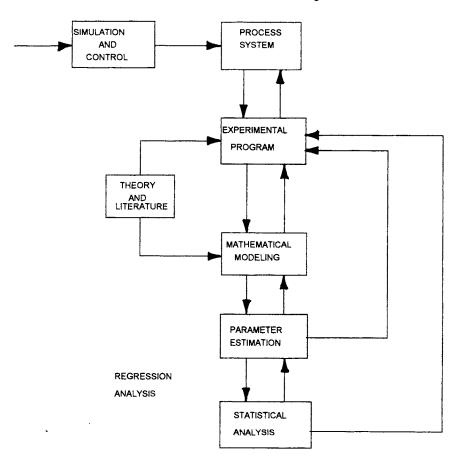


Figure 1-1. Mathematical modeling and regression analysis. By permission, A. Constantinides, *Applied Numerical Methods With Personal Computers*, McGraw-Hill Book Co., 1987.

least-squares curve fitting programs. Also, we will use linear regression analyses to develop statistical relationships involving two or more variables. The most common type of relation is a linear function. By transforming nonlinear functions, many functional relations are made linear.

For the program, we will correlate X-Y data for the following equations:

$$Y = a + bX \tag{1-1}$$

$$Y = a + bX^2 \tag{1-2}$$

$$Y = a + b/X \tag{1-3}$$

$$Y = a + bX^{0.5}$$
 (1-4)

$$Y = aX^{b}$$
(1-5)

$$Y = ae^{bX}$$
(1-6)

$$Y = a + b \log X \tag{1-7}$$

$$Y = a + be^{x} \tag{1-8}$$

We can transform the nonlinear equations (1-5, 1-6, and 1-8) by linearizing as follows:

$Y = aX^{b}$	$\ln Y = \ln a + b \ln X$	(1-9)
$Y = ae^{bX}$	$\ln Y = \ln a + bX$	(1-10)
$Y = a + be^{x}$	$\ln Y = \ln a + (\ln b)X$	(1-11)

LINEAR REGRESSION ANALYSIS

Regression analysis uses statistical and mathematical methods to analyze experimental data and to fit mathematical models to these data. We can solve for the unknown parameters after fitting the model to the data. Suppose we want to find a linear function that involves paired observations on two variables, X the independent variable and Y the dependent variable. Where

 $\begin{array}{l} n = the \ number \ of \ observations \\ X_i = the \ i^{th} \ observation \ of \ the \ independent \ variable \\ Y_i = the \ i^{th} \ observation \ of \ the \ dependent \ variable \end{array}$

We can develop a linear regression model that expresses Y as a function of X. We can further derive formulae to determine the values of a and b that give the best fit of the equations. For each experimental point corresponding to an X-Y pair, there will be an element that represents the difference between the corresponding calculated value \hat{Y} and the original value of Y. This is expressed as:

$$\mathbf{r}_{i} = \hat{\mathbf{Y}}_{i} - \mathbf{Y} \tag{1-12}$$

 r_i may be either positive or negative depending on the side of the fitted curve of X-Y points. Here, we can minimize the sum of the squares of the residuals by the following expression:

SRS =
$$\sum_{i=1}^{n} r_i^2 = \sum_{i=1}^{n} \zeta_i^2 = \text{minimum}$$
 (1-13)

where

n is the number of observations of X-Y points

$$\hat{\mathbf{Y}}_{i} = \mathbf{a} + \mathbf{b}\mathbf{X}_{i} \tag{1-14}$$

$$\mathbf{r}_i = \mathbf{a} + \mathbf{b}\mathbf{X}_i - \mathbf{Y}_i \tag{1-15}$$

and

SRS =
$$\sum_{i=1}^{n} r_i^2 = \sum_{i=1}^{n} (a + bX_i - Y_i)^2$$
 (1-16)

The problem is reduced to finding the values of a and b so that the summation of Equation 1-16 is minimized. We can obtain this by taking the partial derivative of Equation 1-16 with respect to each variable a and b and set the result to zero.

$$\frac{\partial \sum r_i^2}{\partial a} = 0 \quad \text{and} \quad \frac{\partial \sum r_i^2}{\partial b} = 0 \quad (1-17)$$

Substituting Equation 1-16 into Equation 1-17, we obtain

$$\frac{\partial \sum (a + bX_i - Y_i)^2}{\partial a} = 0$$
(1-18)

and

$$\frac{\partial \sum (a + bX_i - Y_i)^2}{\partial b} = 0$$
(1-19)

This is equivalent to

$$\frac{2\sum (a + bX_i - Y_i)\partial \sum (a + bX_i - Y_i)}{\partial a} = 0$$
(1-20)

and

$$(1-21)$$

Since b, X, and Y are not functions of a, and the partial derivative of a with respect to itself is unity, Equation 1-20 reduces to

$$\sum a + \sum bX_i = \sum Y_i \tag{1-22}$$

Similarly, a, X, and Y are not functions of b. Therefore, Equation 1-21 becomes

$$\sum aX_i + \sum bX_i^2 = \sum X_i Y_i$$
(1-23)

where a and b are constants. Equations 1-18 and 1-19 are expressed as:

$$an + b\sum X_i = \sum Y_i \tag{1-24}$$

and

$$a\sum X_{i} + b\sum X_{i}^{2} = \sum X_{i}Y_{i}$$
(1-25)

We have now reduced the problem of finding a straight line through a set of X - Y data points to one of solving two simultaneous equations 1-24 and 1-25. Both equations are linear in X, Y, and n, and the unknowns a and b. Using Cramer's rule for the simultaneous equations, we have

$$a = \frac{\begin{vmatrix} \sum Y_{i} & \sum X_{i} \\ \sum X_{i}Y_{i} & \sum X_{i} \end{vmatrix}}{\begin{vmatrix} n & \sum X_{i} \\ \sum X_{i} & \sum X_{i} \end{vmatrix}}$$
(1-26)

and

$$b = \frac{\begin{vmatrix} n & \sum Y_i \\ \sum X_i & \sum X_i Y_i \end{vmatrix}}{\begin{vmatrix} n & \sum X_i \\ \sum X_i & \sum X_i^2 \end{vmatrix}}$$
(1-27)

Solving Equations 1-26 and 1-27 gives

$$a = \frac{\sum X_{i}^{2} \sum Y_{i} - \sum X_{i} \sum X_{i} Y_{i}}{n \sum X_{i}^{2} - \sum X_{i} \sum X_{i}}$$
(1-28)

and

$$b = \frac{n \sum X_i Y_i - \sum X_i \sum Y_i}{n \sum X_i^2 - \sum X_i \sum X_i}$$
(1-29)

respectively, where all the sums are taken over all experimental observations.

Also, we can construct a table with columns headed X_i , Y_i , X_i^2 , X_iY_i as an alternative to obtain the values of a and b. The sums of the columns will then give all the values required to Equations 1-28 and 1-29.

METHODS FOR MEASURING REGRESSION

Linear Regression

From the values of a, b, and X_i , the independent variable, we can now calculate the corresponding estimated value of Y, designated as \hat{Y}_i .

$$\hat{\mathbf{Y}}_{i} = \mathbf{a} + \mathbf{b} \cdot \mathbf{X}_{i} \tag{1-30}$$

Figure 1-2 illustrates this equation known as the regression line. The variation of the observed Y's about the mean of the observed Y's is the total sum of squares, and can be expressed as:

$$SST = \sum (Y_i - \overline{Y})^2$$
(1-31)

where

$$\overline{Y} = \frac{\sum Y_i}{N}$$
(1-32)

Figure 1-3 shows the differences between Y and \overline{Y} , which is the total sum of squares. This is the sum of these squared differences. The total sum of squares can be expressed as:

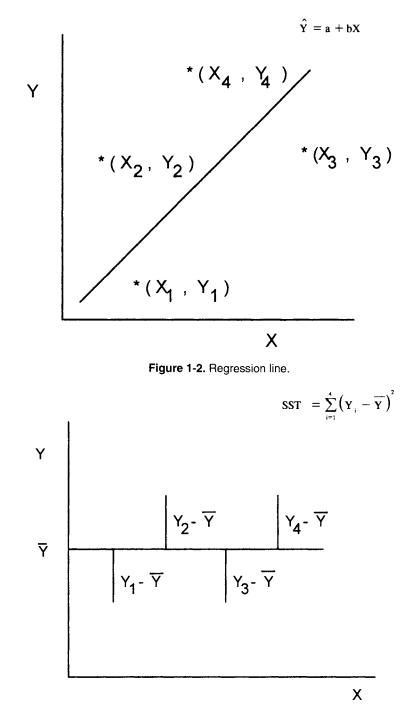


Figure 1-3. Total sum of squares.

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$$SST = \sum (Y_i - \overline{Y})^2$$
(1-33)

$$= \sum \left[(\mathbf{Y}_{i} - \hat{\mathbf{Y}}_{i}) + (\hat{\mathbf{Y}}_{i} - \overline{\mathbf{Y}}) \right]^{2}$$
(1-34)

$$= \sum (Y_{i} - \hat{Y}_{i})^{2} + 2\sum (Y_{i} - \hat{Y}_{i})(\hat{Y}_{i} - \overline{Y}) + \sum (\hat{Y}_{i} - \overline{Y})^{2}$$
(1-35)

and since

$$2\sum (\mathbf{Y}_{i} - \hat{\mathbf{Y}}_{i})(\hat{\mathbf{Y}}_{i} - \overline{\mathbf{Y}}) = 0$$

then

$$SST = \sum (Y_i - \hat{Y}_i)^2 + \sum (\hat{Y}_i - \overline{Y})^2$$

$$= SSE + SSR$$
(1-36)

where

$$SSE = \sum (Y_{i} - \hat{Y}_{i})^{2}$$
(1-37)

and

$$SSR = \sum (\hat{Y}_i - \overline{Y})^2$$
(1-38)

SSE is the error (or residual) sum of squares. This is the quantity represented by Equation 1-13, that is, the equation we want to minimize. It is the sum of the differences squared between the observed Y's and the estimated (or computed) \hat{Y} 's. Figure 1-4 shows the differences between the Y's and \hat{Y} 's. SSR is the regression sum of squares and measures the variation of the estimated values, \hat{Y} , about the mean of the observed Y's and \overline{Y} . Figure 1-5 shows the differences between Y's and \overline{Y} . The regression sum of squares is the sum of the squared differences.

The ratio of the regression sum of squares to the total sum of squares is used in calculating the coefficient of determination. This shows how well a regression line fits the observed data. The coefficient of determination is:

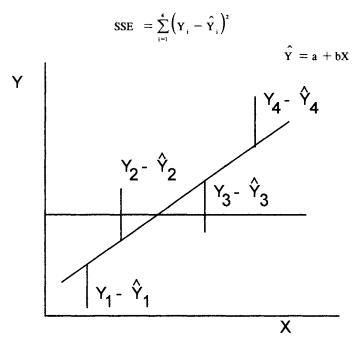


Figure 1-4. Error sum of squares.

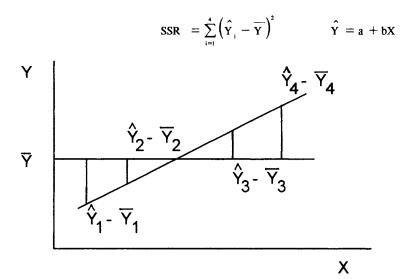


Figure 1-5. Regression sum of squares.

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$$r^2 = \frac{SSR}{SST}$$
(1-39)

and since SSR = SST - SSE

$$r^2 = 1 - \frac{SSE}{SST}$$
(1-40)

The coefficient of determination has the following properties:

- 1. $0 \le r^2 \le 1$
- 2. If $r^2 = 1$ all ξ_i are zero. The observed Y's and the estimated \hat{Y} 's are the same and this indicates a perfect fit.
- 3. If $r^2 = 0$, no linear functional relationship exists.
- 4. As r² approaches one, the better the fit. The closer r² approaches zero, the worse the fit. The correlation coefficient r is:

$$r = \left(1 - \frac{SSE}{SST}\right)^{0.5}$$
(1-41)

Although the correlation coefficient gives a measure of the accuracy of fit, we should treat this method of analysis with great caution. Because r is close to one does not always mean that the fit is necessarily good. It is possible to obtain a high value of r when the underlying relationship between X and Y is not even linear. Draper and Smith [6] provide an excellent guidance of assessing results of linear regression.

THE ANALYSIS OF VARIANCE TABLE FOR LINEAR REGRESSION

Each sum of squares SSR, SSE, and SST corresponds to many degrees of freedom. This is the number of observations, n minus the number of independent quantities. The error sum of squares SSE has N-2 degrees of freedom since

$$SSE = \sum (Y_i - \hat{Y}_i)^2$$

and

$$\hat{\mathbf{Y}} = \mathbf{a} + \mathbf{b} \cdot \mathbf{X}$$

then

SSE =
$$\sum [Y_i - (a + bX_i)]^2$$
 (1-42)

where a and b represent two independent quantities calculated from a set of N observations giving the error sum of squares N-2 degrees of freedom.

The error mean squares, MSE, is an estimate of the variance of the error of observation, ξ , and is:

$$MSE = \frac{SSE}{N-2}$$
(1-43)

The number of degrees of freedom for the total sums of squares SST is N-1 since

$$SST = \sum (Y_i - \overline{Y}_i)^2$$

 \overline{Y} represents a quantity from a set of N observations. This gives the total sum of squares N-1 degrees of freedom. The total mean squares, MST, is an estimate of the variance of the dependent variable Y and is:

$$MST = \frac{SST}{N-1}$$
(1-44)

The regression sum of squares, SSR, has one degree of freedom. Since

$$SSR = SST - SSE$$

= (N - 1) - (N - 2)
= 1

The regression mean squares, MSR, is an estimate of the variance of the estimates, \hat{Y} , and since the regression sum of squares has one degree of freedom, the regression mean squares is

$$MSR = SSR \tag{1-45}$$

Table 1-1 shows the variance table for linear regression.

The test statistics for linear regression is:

$$F = \frac{MSR}{MSE}$$
(1-46)

Source of Variation	Degrees of Freedom	Sum of Squares	Mean Squares
Total	N - 1	$SST = \sum (Y_i - \overline{Y})^2$	$MST = \frac{SST}{N-1}$
Regression	1	$SSR = \sum (\hat{Y} - \overline{Y})^2$	MSR = SSR
Error	N-2	$SSE = \sum (Y_i - \hat{Y}_i)^2$	$MSE = \frac{SSE}{N-2}$

Table 1-1Variance Table for Linear Regression

The F-distribution has one degree of freedom in the numerator and N-2 degree of freedom in the denominator.

MULTIPLE REGRESSION ANALYSIS

Inadequate results are sometimes obtained with a single independent variable. This shows that one independent variable does not provide enough information to predict the corresponding value of the dependent variable. We can approach this problem, if we use additional independent variables and develop a multiple regression analysis to achieve a meaningful relationship. Here, we can employ a linear regression model in cases where the dependent variable is affected by two or more controlled variables.

The linear multiple regression equation is expressed as:

$$Y = C_0 + C_1 X_1 + C_2 X_2 + \dots + C_K X_K$$
(1-47)

where Y = the dependent variable $X_1, X_2, \dots X_K =$ the independent variables K = the number of independent variables $C_0, C_1, C_2, \dots C_K =$ the unknown regression coefficients

The unknown coefficients are estimated based on n observation for the dependent variable Y, and for each of the independent variables X_i 's where i = 1, 2, 3, ..., K.

These observations are of the form:

$$Y_{j} = C_{0} = C_{1}X_{1j} + C_{2}X_{2j} + \dots + C_{K}X_{Kj} + \xi_{j}$$
(1-48)

for j = 1, 2, ... N

where $Y_j = \text{the } j_{th}$ observation of the dependent variable $X_{1j}, \ldots X_{Kj} = \text{the } j_{th}$ observation of the $X_1, X_2, \ldots X_K$ independent variables

We can use a least squares technique to calculate estimates of $\hat{C}_0, \hat{C}_1, \dots \hat{C}_K$ of the coefficients $C_0, C_1, \dots C_K$ by minimizing the following equation:

$$S = \sum_{j=1}^{N} \left[Y_{j} - (\hat{C}_{0} + \hat{C}_{1}X_{1j} + \dots + \hat{C}_{K}X_{Kj}) \right]^{2} = \sum_{j=1}^{N} \xi_{j}^{2}$$
(1-49)

Taking the partial derivatives of S with respect to $\hat{C}_0, \hat{C}_1, ..., \hat{C}_K$, that is $\partial s/\partial \hat{C}_0, \partial s/\partial \hat{C}_1, ... \partial s/\partial \hat{C}_K$ and setting them equal to zero, we obtain the following set of equations:

$$\begin{split} &N\hat{C}_{0} + \left(\sum X_{1j}\right)\hat{C}_{1} + \ldots + \left(\sum X_{Kj}\right)\hat{C}_{K} = \sum Y_{j} \\ &\left(\sum X_{1j}\right)\hat{C}_{0} + \left(\sum X_{1j}^{2}\right)\hat{C}_{1} + \ldots + \left(\sum X_{1j}X_{kj}\right)\hat{C}_{K} = \sum X_{1j}Y_{j} \\ &\left(\sum X_{2j}\right)\hat{C}_{0} + \left(\sum X_{1j}X_{2j}\right)\hat{C}_{1} + \ldots + \left(\sum X_{2j}X_{Kj}\right)\hat{C}_{K} = \sum X_{2j}Y_{j} \\ &\vdots \\ &\vdots \\ &\vdots \\ &\vdots \\ &\vdots \\ &\left(\sum X_{Kj}\right)\hat{C}_{0} + \left(\sum X_{1j}X_{Kj}\right)\hat{C}_{1} + \ldots + \left(\sum X_{Kj}^{2}\right)\hat{C}_{K} = \sum X_{Kj}Y_{j} (1-50) \end{split}$$

Equation 1-50 can be expressed in matrix form as:

$$U\hat{C} = V$$

where

$$\mathbf{U} = \begin{bmatrix} \mathbf{N} & \sum X_{1j} & \dots & \dots & \sum X_{Kj} \\ \sum X_{1j} & \sum X_{1j}^2 & \dots & \dots & \sum X_{1j} X_{Kj} \\ \vdots & & & \\ \sum X_{Kj} & \sum X_{1j} X_{Kj} & & \sum X_{Kj}^2 \end{bmatrix}$$

$$\hat{\mathbf{C}} = \begin{bmatrix} \hat{\mathbf{C}}_{0} \\ \hat{\mathbf{C}}_{1} \\ \vdots \\ \vdots \\ \hat{\mathbf{C}}_{K} \end{bmatrix} \qquad \qquad \mathbf{V} = \begin{bmatrix} \sum \mathbf{Y}_{j} \\ \sum \mathbf{X}_{1j} \mathbf{Y}_{j} \\ \vdots \\ \vdots \\ \sum \mathbf{X}_{Kj} \mathbf{Y}_{j} \end{bmatrix}$$

U is a symmetric matrix.

We can obtain estimates for the coefficients $\hat{C}_0, \hat{C}_1, \dots \hat{C}_K$ by successive elimination or by solving for the inverse of U. That is

$$\hat{C} = U^{-1}V$$

where

 U^{-1} = the inverse of U

After solving for $\hat{C}_0, \hat{C}_1, \dots \hat{C}_K$, the estimates of the dependent variable observations \hat{Y}_i can be obtained as follows:

$$\hat{\mathbf{Y}}_{j} = \hat{\mathbf{C}}_{0} = \hat{\mathbf{C}}_{1} \mathbf{X}_{1j} + \dots + \hat{\mathbf{C}}_{K} \mathbf{X}_{Kj}$$
 (1-51)

The power equations have often been derived to calculate the parameters of experimental data. Such an equation can be expressed in the form:

$$Y = C_0 X_1^{C_1} X_2^{C_2} \dots X_K^{C_K}$$
(1-52)

We can calculate the coefficients of the independent variables, if Equation 1-52 is linearized by taking its natural logarithm to give

$$\ln Y = \ln C_0 + C_1 \ln X_1 + C_2 \ln X_2 + \dots + C_K \ln X_K$$
(1-53)

The coefficients $C_0, C_1, C_2, \dots C_K$ can then be obtained by Gaussian elimination. Table 1-2 shows the variance table for linear multiple regression.

The coefficient of determination is

$$r^2 = 1 - \frac{SSE}{SST}$$
(1-54)

and the correlation coefficient is

$$\mathbf{r} = \left(1 - \frac{SSE}{SST}\right)^{0.5} \tag{1-55}$$

The test statistic is the F - ratio, which we can define as:

$$F = \frac{MSR}{MSE}$$
(1-56)

POLYNOMIAL REGRESSION

Some engineering data are often poorly represented by a linear regression. If we know how Y depends on X, then we can develop some form of nonlinear regression [7], although total convergence of this iterative regression procedure can not be guaranteed. However, if the form of dependence is unknown, then we can treat Y as a general function of X by trigonometric terms (Fourier analysis) or polynomial function. The

 Table 1-2

 Analysis of Variance Table for Linear Multiple Regression

Source of Variance	Degree of Freedom	Sum of Squares	Mean Squares
Total	N – 1	$SST = \sum (Y_j - \overline{Y})^2$	$MST = \frac{SST}{N-1}$
Regression	К	$SSR = \sum (\hat{Y}_j - \overline{Y})^2$	$MSR = \frac{SSR}{K}$
Error	N – K – 1	$SSE = \sum (Y_j - \hat{Y}_j)^2$	$MSE = \frac{SSE}{N - K - 1}$

least squares procedure can be readily extended to fit the data to an nth-degree polynomial:

$$Y = C_0 + C_1 X + C_2 X^2 + \dots C_n X^n$$
 (1-57)

where $C_0, C_1, C_2, \ldots C_n$ are constants.

For this case, the sum of the squares of the residuals is minimized:

$$S = \sum_{j=1}^{N} \left[Y_{j} - C_{0} - C_{1}X_{j} - C_{2}X_{j}^{2} - \dots - C_{n}X_{j}^{n} \right]^{2}$$
(1-58)

At the minimum, all the partial derivatives with respect to the chosen constants are zero; i.e.,

$$\frac{\partial S}{\partial C_0}, \frac{\partial S}{\partial C_1}, \dots, \frac{\partial S}{\partial C_n} = 0$$

This gives a system of (n+1) linear equations in (n+1) unknowns, C_0 , C_1 , ..., C_n

$$\frac{\partial S}{\partial C_0} = 0 = \sum_{j=1}^{N} 2(Y_j - C_0 - C_1 X_j - C_2 X_j^2 - \dots - C_n X_j^n)(-1)$$

$$\frac{\partial S}{\partial C_1} = 0 = \sum_{j=1}^{N} 2(Y_j - C_0 - C_1 X_j - C_2 X_j^2 - \dots - C_n X_j^n)(-X_j)$$

$$\frac{\partial S}{\partial C_2} = 0 = \sum_{j=1}^{N} 2(Y_j - C_0 - C_1 X_j - C_2 X_j^2 - \dots - C_n X_j^n)(-X_j^2)$$

$$\vdots$$

$$\vdots$$

$$\vdots$$

$$\frac{\partial S}{\partial C_n} = 0 = \sum_{j=1}^{N} 2(Y_j - C_0 - C_1 X_j - C_2 X_j^2 - \dots - C_n X_j^n)(-X_j^n)$$

The above equations are set to equal zero and can be rearranged in the following set of normal equations.

$$C_{0}N + C_{1}\sum X_{j} + C_{2}\sum X_{j}^{2} + \dots + C_{n}\sum X_{j}^{n} = \sum Y_{j}$$

$$C_{0}\sum X_{j} + C_{1}\sum X_{j}^{2} + C_{2}\sum X_{j}^{3} + \dots + C_{n}\sum X_{j}^{n+1} = \sum X_{j}Y_{j}$$

$$C_{0}\sum X_{j}^{2} + C_{1}\sum X_{j}^{3} + C_{2}\sum X_{j}^{4} + \dots + C_{n}\sum X_{j}^{n+2} = \sum X_{j}^{2}Y_{j}$$

$$\vdots$$

$$\vdots$$

$$C_{0}\sum X_{j}^{n} + C_{1}\sum X_{j}^{n+1} + C_{2}\sum X_{j}^{n+2} + \dots + C_{n}\sum X_{j}^{2n} = \sum X_{j}^{n}Y_{j}$$
(1-60)

Equation 1-60 in matrix form, becomes

UC=V

$$U = \begin{bmatrix} N & \sum X_{j} & \sum X_{j}^{2} & \dots & \sum X_{j}^{n} \\ \sum X_{j} & \sum X_{j}^{2} & \sum X_{j}^{3} & \dots & \sum X_{j}^{n+1} \\ \sum X_{j}^{2} & \sum X_{j}^{3} & \sum X_{j}^{4} & \dots & \sum X_{j}^{n+2} \\ \vdots & \vdots & \vdots & \vdots & \vdots \\ \sum X_{j}^{n} & \sum X_{j}^{n+1} & \sum X_{j}^{n+2} & \dots & \sum X_{j}^{2n} \end{bmatrix}$$
$$C = \begin{bmatrix} C_{0} \\ C_{1} \\ C_{2} \\ \vdots \\ C_{n} \end{bmatrix} \qquad V = \begin{bmatrix} \sum Y_{j} \\ \sum X_{j}^{2} Y_{j} \\ \sum X_{j}^{2} Y_{j} \\ \vdots \\ \sum X_{j}^{n} Y_{j} \end{bmatrix}$$

Linear equations generated by polynomial regression can be ill-conditioned when the coefficients have very small and very large numbers. This results in smooth curves that fit poorly. Ill-conditioning often happens if the degree of the polynomial is large and if the Y values cover a wide range. We can determine the error of polynomial regression by a standard error of the estimate as

$$\sigma^2 = \frac{\sum SSE}{N - n - 1} \tag{1-61}$$

where

n = the degree of polynomial N = the number of data pairs

The coefficient of determination can be expressed as:

$$r^{2} = 1 - \frac{SSE}{SST}$$

= $1 - \frac{\sum_{j=1}^{N} (Y_{j} - \hat{Y}_{j})^{2}}{\sum_{j=1}^{N} (Y_{j} - \overline{Y}_{j})^{2}}$ (1-62)

The correlation coefficient is given by

$$\mathbf{r} = \left(1 - \frac{SSE}{SST}\right)^{0.5} \tag{1-63}$$

The numerator of Equation 1-61 should continually decrease as the degree of the polynomial is raised. Alternatively, the denominator of Equation 1-61 causes σ^2 to increase as we move away from the optimum degree.

SOLUTION OF SIMULTANEOUS LINEAR EQUATIONS

Analyses of physiochemical systems often give us a set of linear algebraic equations. Also, methods of solution of differential equations and nonlinear equations use the technique of linearizing the models. This requires repetitive solutions of sets of linear algebraic equations. Linear equations can vary from a set of two to a set having 100 or more equations. In most cases, we can employ Cramer's rule to solve a set of two or three linear algebraic equations. However, for systems of many linear equations, the algebraic computation becomes too complex and may require other methods of analysis.

We will analyze two methods of solving a set of linear algebraic equations, namely Gauss elimination and Gauss-Seidel iteration methods. Gauss elimination is most widely used to solve a set of linear algebraic equations. Other methods of solving linear equations are Gauss-Jordan and LU decomposition. Table 1-3 illustrates the main advantages and disadvantages of using Gauss, Gauss-Jordan and LU decomposition.

Gaussian elimination method is based upon the principle of converting a set of N equations of N unknowns represented by

$$a_{1,1}X_{1} + a_{1,2}X_{2} + a_{1,3}X_{3} + \dots + a_{1,N}X_{N} = Y_{1}$$

$$a_{2,1}X_{1} + a_{2,2}X_{2} + a_{2,3}X_{3} + \dots + a_{2,N}X_{N} = Y_{2}$$

$$\vdots$$

$$a_{N,1}X_{1} + a_{N,2}X_{2} + a_{N,3}X_{3} + \dots + a_{N,N}X_{N} = Y_{N}$$
(1-64)

•		·
Method	Advantages	Disadvantages
Gauss elimination	The most fundamental solution algorithm.	Solution of one set of linear equations at a time.
Gauss-Jordan	Basis for computing inverse; can solve multiple sets of equations.	Less efficient for a single set of equations.
LU decomposition	Efficient if one set of linear equations is repeatedly solved with different inhomogeneous terms (e.g., in the inverse power method.)	Less efficient and more cumbersome than Gauss elimination if used only once.

 Table 1-3

 Comparison of Three Methods of Linear Equations

Source: S. Nakamura, Applied Numercial Methods With Software, Prentice-Hall Int. Ed., N.J., 1991. [2]

to a triangular set of the form

$$a_{1,1}X_{1} + a_{1,2}X_{2} + a_{1,3}X_{3} + \dots + a_{1,N}X_{N} = Y_{1}$$

$$a'_{2,2}X_{2} + a'_{2,3}X_{3} + \dots + a'_{2,N}X_{N} = Y'_{2}$$

$$a''_{3,3}X_{3} + \dots + a''_{3,N}X_{N} = Y''_{3}$$

$$\vdots$$

$$a^{(N-1)}_{N,N}X_{N} = Y^{(N-1)}_{N} \qquad (1-65)$$

The process involves converting the general form

aX = Y

to the triangular form

$$UX = Y'$$

where U is the upper triangular matrix. After completing the forward elimination process, we then employ the back substitution method starting with the last equation to obtain the solution of the set of the remaining equations. The solution of X_N is obtained from the last equation

$$X_{N} = \frac{Y_{N}^{(N-1)}}{a_{N,N}^{(N-1)}}$$

subsequent solutions are

$$X_{N-1} = \frac{\left[Y_{N-1}^{(N-2)} - a_{N-1,N}^{(N-1)} X_{N}\right]}{a_{N-1,N-1}^{(N-2)}}$$

:
$$X_{1} = \frac{\left[Y_{1} - \sum_{j=2}^{N} a_{1,j} X_{j}\right]}{a_{1,1}}$$

(1-66)

This procedure completes the Gauss elimination. We can carry out the elimination process by writing only the coefficients and the matrix vector in an array as

The array after the forward elimination becomes

We can summarize the operations of Gauss elimination in a form suitable for a computer program as follows:

- 1. Augment the N \times N coefficient matrix with the vector of right hand sides to form a N \times (N-1) matrix.
- 2. Interchange the rows if required such that a_{11} is the largest magnitude of any coefficient in the first column.
- 3. Create zeros in the second through Nth rows in the first column by subtracting a_{i1}/a_{11} times the first row from the /ith row. Store the a_{i1}/a_{11} in a_{i1} , i=2, 3, ... N.
- 4. Repeat Steps 2 and 3 for the second through the $(N-1)^{st}$ rows, putting the largest-magnitude coefficient on the diagonal by interchanging rows (considering only rows j to N). Then subtract a_{ij}/a_{jj} times the jth row from the ith row to create zeros in all the positions of the jth column below the diagonal. Store the a_{ij}/a_{jj} in a_{ij} , $i=j+1 \ldots N$. At the end of this step, the procedure is an upper triangular.
- 5. Solve for X_N from the Nth equation by

$$X_{N} = \frac{a_{N,N+1}}{a_{N,N}}$$

6. Solve for X_{N-1} , X_{N-2} , ..., X_1 from the $(N-1)^{st}$ through the first equation in turn by

$$X_{i} = \frac{a_{i,N+1} - \sum_{j=i+1}^{N} a_{ij} \cdot X_{j}}{a_{ii}}$$

The Guass-Seidel Iterative Method

Some engineering problems give sets of simultaneous linear equations that are diagonally dominant. This involves the computation of finite difference equations, derived from the approximation of partial differential equations. A diagonally dominant system of linear equations has coefficients on the diagonal that are larger in absolute value than the sum of the absolute values of the other coefficients.

Solving a system of N linear equations using the Gauss-Seidel iterative method, we can rearrange the rows so that the diagonal elements have larger absolute value than the sum of the absolute values of the other coefficients in the same row. This is defined as

We start with an initial approximation to the solution vector, $X^{(1)}$, and then calculate each component of $X^{(n+1)}$, for i = 1, 2, 3, ... N by

$$X_{i}^{(n+1)} = \frac{Y_{i}}{a_{II}} - \sum_{j=1}^{i-1} \frac{a_{ij}}{a_{ii}} X_{j}^{(n+1)} - \sum_{j=i+1}^{N} \frac{a_{ij}}{a_{ii}} X_{j}^{n}, n = 1, 2...$$
(1-69)

A necessary condition for convergence is that

$$|a_{ii}| > \sum_{\substack{i=1, \ j \neq i}}^{N} |a_{ij}|$$
 $i = 1, 2, ... N$

When this condition exists, X^N will converge to the correct solution no matter what initial vector is used.

The Gauss-Seidel iterative method requires an initial approximation of the values of the unknowns X_1 to X_N . We use these values in Equation 1-69 to start calculation of new estimates of X's. Each newly calculated X_i replaces its previous value in subsequent calculations. The iteration continues until all the newly calculated X's converge to a convergence criterion, ε , of the previous values. The Gauss-Seidel method is very efficient because of this faster convergence and should be used when the system is diagonally dominant. This method is widely used in the solution of engineering problems. However, the iterative method will not converge for all sets of equations or for all possible rearrangements of the equations. Assume that we are given a set of N simultaneous linear equations:

$$a_{1,1}X_{1} + a_{1,2}X_{2} + \dots + a_{1,N}X_{N} = a_{1,N+1}$$

$$a_{2,1}X_{1} + a_{2,2}X_{2} + \dots + a_{2,N}X_{N} = a_{2,N+1}$$

$$\vdots$$

$$a_{N,1}X_{1} + a_{N,2}X_{2} + \dots + a_{N,N}X_{N} = a_{N,N+1}$$
(1-70)

in which a_{ii}'s are constants.

Method of Solution

We first normalize the coefficients of Equation 1-70 to reduce the number of divisions required in the calculations. This is achieved by dividing all elements in row i by a_{ii} , i = 1, 2, ... N to produce an augmented coefficient matrix given by

$$\begin{bmatrix} 1 & a'_{1,2} & a'_{1,3} & \dots & a'_{1,N} & a'_{1,N+1} \\ a'_{2,1} & 1 & a'_{2,3} & \dots & a'_{2,N} & a'_{2,N+1} \\ \vdots & & & & & \\ a'_{N,1} & a'_{N,2} & a'_{N,3} & \dots & 1 & a'_{N,N+1} \end{bmatrix}$$
(1-71)

where $a'_{i,j} = \frac{a_{i,j}}{a_{i,j}}$

The approximation to the solution vector after the kth iteration is

$$X_{k} = [X_{1,k}, X_{2,k} \dots X_{N,K}]$$

This is modified by the algorithm

$$X_{i,k+1} = a'_{i,N+1} - \sum_{j=1}^{i-1} a'_{ij} \cdot X_{j,K+1} - \sum_{j=i+1}^{N} a'_{ij} \cdot X_{jk}$$
(1-71)

where i = 1, 2, ... N

The next approximation yields

 $\mathbf{X}_{K+1} = [\mathbf{X}_{1,K+1}, \mathbf{X}_{2,K+1} \dots \mathbf{X}_{N,K+1}]$

The iteration subscript K can be omitted because the new values $X_{i,K+1}$ replace the old values during computation and Equation 1-71 becomes

$$X_{i} = a'_{i,N+1} - \sum_{\substack{i=1\\j\neq i}}^{N} a'_{ij}, \quad i = 1, 2, \dots N$$
 (1-72)

The X_i values calculated by iterating with Equation 1-71 will converge to the solution of Equation 1-72, if the convergence criterion is

$$|X_{i,K+1} - X_i| < \varepsilon, \quad i = 1, 2, ... N$$
 (1-73)

provided that no element of the solution vector may have its magnitude changed by an amount greater than ε as a result of one Gauss-Seidel iteration. We can introduce an upper limit on the number of iterations, K_{max} , to terminate the iterative sequence when convergence does not occur.

SOLUTION OF NONLINEAR EQUATIONS

Solving problems in chemical engineering and science often requires finding the real root of a single nonlinear equation. Examples of such computations are in fluid flow, where pressure loss of an incompressible turbulent fluid is evaluated. The Colebrook [8] implicit equation for the Darcy friction factor, f_D , for turbulent flow is expressed

$$\frac{1}{f_{\rm D}^{0.5}} = -2\log\left\{\frac{\varepsilon / D}{3.7} + \frac{2.51}{N_{\rm Re}}f_{\rm D}^{0.5}\right\}$$
(1-74)

where ε/D is the pipe roughness in feet, f_D is the Darcy friction factor, which is four times the Fanning friction factor, and N_{Re} is the Reynolds number. Equation 1-74 is nonlinear and involves trial and error solutions to achieve a solution for f_D ; i.e., the root of the equation.

Equation 1-74 can be further expressed as:

$$F(f_{\rm D}) = \frac{1}{f_{\rm D}^{0.5}} + 2\log\left\{\frac{\epsilon / D}{3.7} + \frac{2.51}{N_{\rm Re}f_{\rm D}^{0.5}}\right\}$$
(1-75)

In the form of natural logarithm, Equation 1-75 is

$$F(f_{\rm D}) = \frac{1}{f_{\rm D}^{0.5}} + 0.86858 \ln\left\{\frac{\varepsilon / D}{3.7} + \frac{2.51}{N_{\rm Re} f_{\rm D}^{0.5}}\right\}$$
(1-76)

The derivative of Equation 1-76 is

$$F'(f_{\rm D}) = -\frac{0.5}{f_{\rm D}^{1.5}} - \frac{4.03329}{\left[N_{\rm Re}f_{\rm D}^{1.5}\frac{\epsilon}{\rm D} + 9.287f_{\rm D}\right]}$$
(1-77)

In thermodynamics, the pressure-volume temperature relationships of real gases are described by the equation of state. The compressibility factor from the reduced pressure and temperature can be rearranged from Redlich and Kwong [9] to two constant state equations in the form

$$Z = 1 + A - \frac{AB}{A + Z} + \frac{A^{2}B}{AZ + Z^{2}}$$
(1-78)

where

A =
$$\frac{0.0867 p_r}{T_r}$$
 and B = $\frac{4.934}{T_r^{1.5}}$

Equation 1-78 is also nonlinear and can be expressed as:

$$F(Z) = Z - 1 - A + \frac{AB}{A + Z} - \frac{A^2B}{AZ + Z^2}$$
(1-79)

The derivative of Equation 1-79 is

$$F'(Z) = 1 - \frac{AB}{(A+Z)^2} + \frac{A^2B(A+2Z)}{(AZ+Z^2)^2}$$
(1-80)

For multicomponent separations, it is often necessary to estimate the minimum reflux ratio of a fractionating column. A method developed for this purpose by Underwood [10] requires the solution of the equation

$$f(\theta) = \sum_{i=1}^{N} \frac{\alpha_{i} x_{i}}{(\alpha_{i} - \theta)} - 1 + q = 0$$
(1-81)

where i = 1, 2, ... Nand N = the number of components in the feed x_i = the mole fraction of component i α_i = the relative volatility of component i q = the thermal condition of the feed

Equation 1-81 is highly nonlinear and must be solved for θ , the Underwood parameter, or the root of the equation. It has a singularity at each value of $\theta = \alpha_i$. If we can determine with some precision the degree of vaporization of the feed and the distillation composition, then we can also obtain a single value of θ by an iterative solution. The value of θ is generally bounded between the relative volatilities of the light and heavy keys.

That is

 $\theta_{\rm LK} < \theta < \theta_{\rm HK}$

Gjumbir and Olujic [11], Tao [12], and Sargent [13] have published articles for solving nonlinear problems. We will discuss some of the methods used in solving nonlinear equations, and employ one of the methods to solve Equation 1-66. Equation 1-81 uses the bisection method to solve for θ in Chapter 7.

The Bisection Method

The method assumes that f(x) is continuous over the interval (x_1,x_2) with $f(x_1)$ and $f(x_2)$ having opposite signs; i.e., $f(x_1).f(x_2) < 0$. There is at least one root of f(x) = 0 in this interval. If $f(x_1) < 0$, then $f(x_2) > 0$. Therefore the interval between these two points can be bisected. If I_1 represents the subinterval (x_2,x_3) at whose end points f(x) takes opposite signs and $x_3 = (x_1 + x_2)/2$, I_1 can be bisected to give an interval I_2 , at whose endpoints f(x) still has opposite signs. We can continue with this process until a root of the equation is found.

The midpoint of interval I_m , is

$$\mathbf{x}_{m+1} = (\mathbf{x}_{m-1}) + \mathbf{x}_{m})/2 \tag{1-82}$$

which is selected as an approximation to the root. The advantages are its simplicity and the number of iterations that can be predicted, because the chosen interval is halved during each iteration.

$$i = \log\left(\frac{x_2 - x_1}{\varepsilon}\right) / \log 2$$
(1-83)

where i is the number of iterations and ε is the tolerance.

Algorithm for the Bisection Method

Determining a root of f(x) = 0, accurate within a specified tolerance value, requires given values of x_1 and x_2 such that $f(x_1)$ and $f(x_2)$ are of opposite signs. i is the iteration index, imax is the maximum number of iterations, ε is the tolerance, and x_3 is the subinterval midpoint.

The final value of x_3 approximates the root. The method may give a false root if f(x) is discontinuous on (x_1, x_2) .

Method of Linear Interpolation (Regula-Falsi Method)

An initial guess, x_1 , is made and function, f_1 , is calculated. We then evaluate the function f_2 at another point, x_2 , such that the two functions have opposite signs; i.e., $f_1f_2 < 0$. We can calculate a new interval value, x_3 , by linear interpolation, and then evaluate the function f_3 .

From similar triangles,

$$\frac{\mathbf{x}_2 - \mathbf{x}_3}{\mathbf{x}_2 - \mathbf{x}_1} = \frac{\mathbf{f}_2}{\mathbf{f}_2 - \mathbf{f}_1}$$
(1-84)

rearranging, we have

$$\mathbf{x}_3 = \mathbf{x}_2 - \frac{\mathbf{f}_2}{\mathbf{f}_2 - \mathbf{f}_1} (\mathbf{x}_2 - \mathbf{x}_1) \tag{1-85}$$

Generalizing, we obtain the recursive algorithm

$$x_{n+1} = x_n - \frac{f_n}{f_n - f_{n-1}} (x_n - x_{n-1})$$
(1-86)

where n = 1, 2, 3, ... N

If $f_n f_{n+1} < 0$, then the righthand interval contains the root, and x_{n+1} becomes the new x_{n-1} for the next iteration. However, if $f_{n-1}f_{n+1} < 0$, the true root of the equation lies between x_{n-1} and x_{n+1} . x_{n+1} then becomes the new x_n for the next iteration.

Algorithm for the Modified Linear Interpolation Method

Step 1. Set SAVE = $f(x_1)$; set $F_1 = f(x_1)$ and $F_2 = f(x_2)$

Step 2. DO WHILE $|\mathbf{x}_1 - \mathbf{x}_2| \ge$ tolerance value 1, or

 $|f(x_3)| \ge$ tolerance value 2

Step 3. $x_3 = x_2 - F_2(x_2-x_1) / (F_2-F_1)$ If $f(x_3)$ of opposite sign to F_1 , then $x_2 = x_3, F_2 = F_3$, If $f(x_3)$ of same sign as SAVE; then $F_1 = F_2/2$ ENDIF ELSE $x_1 = x_3, F_1 = f(x_3)$ If $f(x_3)$ of same sign as SAVE, then $F_2 = F_2/2$ ENDIF ENDIF ENDIF SAVE = $f(x_3)$

The Newton-Raphson Method

The Newton-Raphson method is widely used in finding the root of nonlinear equations. This method uses the derivative of f(x) at x to estimate a new value of the root. The tangent at x is then extended to intersect the x-axis, and the value of x at this intersection is the new estimate of the root. The desired precision is reached by iteration.

Suppose that f(x), f'(x) and f''(x) are continuous and that f(x) and f'(x) do not vanish for the same value of x, if x^r is an approximation to the root of f(x) = 0.

The second mean value theorem can be expressed as:

$$0 = f(x^{r} + h) = f(x^{r}) + hf'(x^{r}) + \frac{h^{2}}{2}f''(x^{r} + O_{2}h)$$
(1-87)

Neglecting higher order terms, we have

$$h = -\frac{f(x^{r})}{f'(x^{r})}$$
(1-88)

Therefore, the next approximation to the root is given by

$$\mathbf{x}^{r+1} = \mathbf{x}^{r} - \frac{\mathbf{f}(\mathbf{x}^{r})}{\mathbf{f}'(\mathbf{x}^{r})}$$
(1-89)

The procedure is repeated until $|\mathbf{x}^r| < \varepsilon$, where ε is a preset tolerance.

This method converges much faster than the bisection technique. However, its rate of convergence does not necessarily correlate with the nearness of the initial starting of the root. The main advantages of this technique are that it converges rapidly (i.e., quadratic convergence) and needs only a single starting point. Its disadvantages are sensitivity to local maxima and minima and a potentially narrow convergence range.

Algorithm for the Newton-Raphson Method

Step 1. Set i = 1 Step 2. $x_i = -f(x_i)/f'(x_i)$ Step 3. If $|x_{i+1} - x_i| < \varepsilon$, then print root x_{i+1} , quit Step 4. i = i+1 If i > imax, then quit, else repeat Steps 2-4

The Secant Method

In cases where the first derivative cannot easily be determined by the Newton-Raphson method, a simple method to approximate the first derivative can be used. The secant method can be determined from two prior estimates.

$$f'(x_i) = \frac{f(x_i) - f(x_{i-1})}{x_i - x_{i-1}}$$
(1-90)

and the iteration process becomes

$$\mathbf{x}_{i+1} = \mathbf{x}_{i} + f(\mathbf{x}_{i+1}) \frac{\mathbf{x}_{i} - \mathbf{x}_{i-1}}{f(\mathbf{x}_{i}) - f(\mathbf{x}_{i-1})}$$
(1-91)

The advantages of this method are that it offers rapid convergence without requiring the first derivative. Convergence is between linear and quadratic, i.e., a power ranging from 1 to 2. The value is an approximation of the tangent of the secant. The value depends upon the steepness of the curve. Because the denominator always approaches zero near the root, this method is prone to instability. It also requires two initial estimates to start.

INTERPOLATIONS

Experimental and physical property data sometimes require values of their unknown functions that correspond to certain values of their independent variables. In certain cases, we may want to determine the behavior of the function. Alternatively, we may want to approximate other values of the function at values of the independent variables that are not tabulated. We can achieve these objectives either by interpolation or extrapolation of a polynomial that fits a selected set of points of both variables (x_i , $f(x_i)$). Also, we can assume after finding a polynomial that fits a selected set of points (x_i , $f(x_i)$ that the polynomial and the function behave over a given interval in question. The values of the polynomial will be estimates of the values of the unknown function.

Generally, the experimental data are approximated by a polynomial, the degree of which can often be calculated by constructing a difference table. The difference column that gives approximate constant value shows the degree of the polynomial that can be fitted to the data. When the polynomial is of the first degree, we have a linear interpolation. For polynomials of higher degrees, we can approximate functions if we construct a table with wider spacing. Such a table is known as a difference table.

A Difference Table

We can obtain a table of values, if the dependent variable f(x) is a function of the independent variable X. If we let h be the uniform difference in the x-values, $h = \Delta x$, we can define the first differences of the function as:

$$\Delta f(x_{1}) = f(x_{2}) - f(x_{1})$$

$$\Delta f(x_{2}) = f(x_{3}) - f(x_{2})$$
:
$$\Delta f(x_{1}) = f(x_{1+1}) - f(x_{1})$$
(1-92)

or

$$\Delta \mathbf{f}_{i} = \mathbf{f}_{i+1} - \mathbf{f}_{i} \tag{1-93}$$

The second difference is the difference of the first differences and can be expressed as:

$$\Delta^{2} f(\mathbf{x}) = \Delta (\Delta f(\mathbf{x}_{1}))$$
(1-94)
= $\Delta (\Delta f(\mathbf{x}_{2}) - \Delta f(\mathbf{x}_{1}))$
= $(f(\mathbf{x}_{3}) - f(\mathbf{x}_{2})) - (f(\mathbf{x}_{2}) - f(\mathbf{x}_{1}))$ (1-95)

or

$$\Delta^{2} f_{1} = (f_{3} - f_{2}) - (f_{2} - f_{1})$$

= f_{3} - 2f_{2} + f_{1} (1-96)

$$\Delta^2 f_i = f_{i+2} - 2f_{i+1} + f_i$$
(1-97)

$$\Delta^{3} f_{1} = \Delta (\Delta^{2} f_{2} - \Delta^{2} f_{1})$$

= $(f_{4} - 2f_{3} + f_{2}) - (f_{3} - 2f_{2} + f_{1})$
= $f_{4} - 2f_{3} + f_{2} - f_{3} + 2f_{2} - f_{1}$ (1-98)

$$\Delta^3 f_1 = f_4 - 3f_3 + 3f_2 - f_1 \tag{1-99}$$

$$\Delta^3 f_i = f_{i+3} - 3f_{i+2} + 3f_{i+1} - f_i$$
(1-100)

We then have a general formula given by

$$\Delta^{n} f_{i} = f_{i+1} - {n \choose 1} f_{i+n-1} + {n \choose 2} f_{i+n-2} - {n \choose 3} f_{i+n-3} + \dots$$

$$= f_{i+1} - n f_{i+n-1} + \frac{n!}{2!(n-2)!} f_{i+n-2} - \frac{n!}{3!(n-3)!} f_{i+n-3} + \dots$$

$$= f_{i+1} - n f_{i+n-1} + \frac{n(n-1)}{2!} f_{i+n-2} - \frac{n(n-1)(n-2)}{3!} f_{i+n-3} + \dots$$
(1-101)

Equation 1-101 is the array of coefficients in the binomial expansion.

Table 1-4 shows how the differences in the dependent variables are arranged.

Interpolating Polynomials

When the tabulated function resembles a polynomial of a constant value in the nth order differences, we can approximate the function by constructing a polynomial. Various formulas have been expressed for the nth degree polynomial that passes through n + 1 pairs of points (x_i, f_i) , where i = 1, 2 ... n + 1.

When the data points are available at equal intervals of the independent variable, we can use the Newton-Gregory forward polynomial.

Table 1-4Diagonal Difference Table					
x	f(x)	Δf	$\Delta^2 f$	$\Delta^3 f$	$\Delta^4 f$
\mathbf{x}_0	\mathbf{f}_0	• 6			
x ₁	f_1	Δf_0	$\Delta^2 f_0$		
x ₂	\mathbf{f}_2	$\Delta \mathbf{f}_{_{1}}$	$\Delta^2 \mathbf{f}_1$	$\Delta^3 { m f}_0$	$\Delta^4 f_0$
X ₃	f_3	Δf_2	$\Delta^2 f_2$	$\Delta^3 f_1$	
<u>x</u> 4	f_4	Δf_3			

$$P_{n}(x_{s}) = f_{0} + {\binom{s}{1}} \Delta f_{0} + {\binom{s}{2}} \Delta^{2} f_{0} + {\binom{s}{3}} \Delta^{3} f_{0}$$

$$+ {\binom{s}{4}} \Delta^{4} f_{0} + \dots + {\binom{s}{n}} \Delta^{n} f_{0}$$

$$= f_{0} + s \Delta f_{0} + \frac{s(s-1)}{2!} \Delta^{2} f_{0} + \frac{s(s-1)(s-2)}{3!} \Delta^{3} f_{0}$$

$$+ \frac{s(s-1)(s-2)(s-3)}{4!} \Delta^{4} f_{0}$$

$$+ \dots + \frac{1}{n!} s(s-1)(s-2) \dots (s-n+1) \qquad (1-102)$$

where $x = x_0 + sh$ and $h = x_{i+1} - x_i = x_2 - x_1$ The Newton-Gregory backward polynomial can be expressed as:

$$P_{n}(x_{s}) = f_{0} + {s \choose 1} \nabla f_{0} + {s+1 \choose 2} \nabla^{2} f_{0} + {s+2 \choose 3} \nabla^{3} f_{0}$$

$$+ {s+3 \choose 4} \nabla^{4} f_{0} + \dots + {s+n-1 \choose n} \nabla^{n} f_{i}$$

$$= f_{0} + s \nabla f_{0} + \frac{s(s+1)}{2!} \nabla^{2} f_{0} + \frac{s(s+1)(s+2)}{3!} \nabla^{3} f_{0}$$

$$+ \frac{s(s+1)(s+2)(s+3)}{4!} \Delta^{4} f_{0} + \dots$$

$$+ \frac{1}{n!} (s+n-1)(s+n-2) \dots s(s+1) \nabla^{n} f_{i} \qquad (1-103)$$

where s is a local coordinate defined by

$$s = \frac{x - x_i}{h}$$

$$\binom{s+n-1}{n}$$
 is a binomial coefficient and $\nabla^n f_i$ is the backward difference.

Stirling central differences represent a horizontal line using averages of the differences; i.e.,

$$\delta f_{i} = f_{i+1/2} - f_{i-1/2}$$

or

$$\delta f_{i+1/2} = f_{i+1} - f_i$$

where

$$f_{i+1/2} = f\left(x_i + \frac{h}{2}\right)$$

$$p_{n}(x) = f_{0} + {\binom{s}{1}} \frac{\Delta f_{-1} + \Delta f_{0}}{2} + \frac{{\binom{s+1}{2}} + {\binom{s}{2}} \Delta^{2} f_{-1}}{2} + {\binom{s+1}{3}} \frac{\Delta^{3} f_{-2f} + \Delta^{3} f_{-1}}{2} + \frac{{\binom{s+2}{4}} + {\binom{s+1}{4}}}{2} \Delta^{4} f_{-2} + (1-104)$$

The forward and the backward difference approximations give the derivatives of the Newton interpolation polynomial at the edges of the interpolation range. However, the central difference is derived from the Newton interpolation at the center of the range of interpolation. Accuracy of an interpolation formula based on equispaced points is highest at the center of the interpolation range. Therefore, the central difference interpolation formula is always more accurate than the forward or backward difference approximations.

SOLUTION OF INTEGRATION

We frequently use numerical techniques to integrate a function given in both analytical and tabular forms. For instance, we can use an integral method to determine the volumetric rate of a gas through a duct from the linear velocity distribution. In fluid mixing with residence time distribution theory, Danckwerts [14] showed that the fraction of material in the outlet stream that has been in the system for a period between t and t+dt is equal to Edt. E is a function of t, and E(t) is the residence time distribution function. We can express E(t) in integral form as:

$$\int_{0}^{\infty} E(t)dt = 1 \tag{1-105}$$

The average time spent by material flowing at a rate, q, through a volume, V, equals V/q.

The mean residence time,
$$\bar{t} = \frac{V}{q}$$

We can also express Equation 1-105 in the form of dimensionless time where $\theta = tq/V$, and this becomes:

$$\int_{0}^{\infty} E(\theta) d\theta = 1$$
 (1-106)

Figure 1-6 shows a residence time distribution from a tracer experiment studying the mixing characteristics of a nozzle-type reactor [15] that behaves nonideally.

Numerical integral methods can be used to calculate areas under this or similar distribution functions.

The Trapezoidal Rule

This is a numerical integration method derived by integrating the linear interpolation formula. It is expressed as:



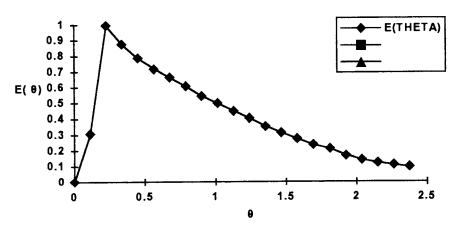


Figure 1-6. Residence time distribution in a nozzle type reactor. Source: A. K. Coker, *A Study of Fast Reactions in Nozzle Type Reactors,* Ph.D. thesis, Univ. of Aston, Birmingham, U.K., 1985.

$$I = \int_{x_{i}}^{x_{i+1}} f(x) dx = \frac{f(x_{i}) + f(x_{i+1})}{2} (\Delta x)$$

$$= \frac{h}{2} (f_{i} + f_{i+1})$$
(1-106)

For (a, b) subdivided into sub-intervals of size h, we can express the area as:

$$I = \int_{a}^{b} f(x) dx = \sum_{i=1}^{n} \frac{h}{2} (f_{i} + f_{i+1})$$

= $\frac{h}{2} (f_{1} + 2f_{2} + 2f_{3} + ... + 2f_{n} + f_{n+1}) + E$ (1-107)

= width \times average height

E represents the local error, $E \cong -\left(\frac{b-a}{12}\right)h^2 f''(\varepsilon)$ where $a \le \varepsilon \le b$

Simpson's 1/3 Rule

Simpson's 1/3 rule is based on quadratic polynomial interpolation. For a quadratic integrated over two Δx intervals that are of uniform width or panels, we can express the area as:

$$I = \int_{a}^{b} f(x) dx = \frac{h}{3} (f_{1} + 4f_{2} + 2f_{3} + 4f_{4} + 2f_{5} + \dots + 2f_{n-1} + 4f_{n} + f_{n+1}) + E$$
(1-108)

= width \times average height

$$E \cong -\frac{(b-a)}{180} h^4 f^{iv}(\epsilon), \quad a \le \epsilon \le b$$

Simpson's 3/8 Rule

Simpson's 3/8 rule is derived by integrating a third-order polynomial interpolation formula. For a domain (a,b) divided into three intervals, it is expressed as:

$$I = \int_{a}^{b} f(x)dx = \frac{3h}{8}(f_{1} + 3f_{2} + 3f_{3} + 2f_{4} + 3f_{5} + 3f_{6} + \dots + 2f_{n-2} + 3f_{n-1} + 3f_{n} + f_{n+1}) + E$$
(1-109)

= width \times average height

$$E \cong -\frac{(b-a)}{80} h^4 f^{i\nu}(\epsilon), \quad a \le \epsilon \le b$$

Gaussian Quadrature

Gauss quadratures are numerical integration methods that employ Legendre points. Gauss quadrature cannot integrate a function given in a tabular form with equispaced intervals. It is expressed as:

$$I = \int_{-1}^{1} f(x)dx = af(x_1) + bf(x_2) + E$$
 (1-110)

where the limits of integration are a to b. To use the tabulated Gaussian quadrature parameters, we must change the interval of integration to (-1,1).

If we let

$$t = \frac{(b-a)x + b + a}{2}$$
 and $dt = \left(\frac{b-a}{2}\right)dx$

then

$$\int_{a}^{b} f(t)dt = \int_{-1}^{1} f(t) \left(\frac{dt}{dx}\right) dx$$
$$= \left(\frac{b-a}{2}\right) \int_{-1}^{1} f\left\{\frac{(b-a)x+b+a}{2}\right\} dx$$
(1-111)

Table 1-5 summarizes the advantages and disadvantages of numerical integration techniques.

SOLUTION OF ORDINARY DIFFERENTIAL EQUATIONS

In certain cases, we may need to find out the behavior of many dynamic and physical processes. This is often expressed mathematically

Method	Advantages	Disadvantages Needs a large number of subintervals for good accuracy.	
Trapezoidal rule	Simplicity. Optimal for improper integrals.		
Simpson's 1/3 rule	Simplicity. Higher accuracy than the trapezoidal rule.	Even number of inter- vals only.	
Simpson's 3/8 rule	Same order of accuracy as the 1/3 rule.	Intervals or panels in multiples of three only.	
Gaussian quadrature	Functional data at two end points are not used.	Data points are not equispaced.	

Table 1-5Advantages and Disadvantages of NumericalIntegration Techniques

by ordinary differential equations. The solutions of these equations are of great value to engineers and scientists. Many differential equations can be solved by well-known analytical methods, although many physically differential equations are impossible to solve analytically. Numerical techniques can be developed to solve these equations. We shall consider the numerical techniques for solving differential equations.

Nth Order Ordinary Differential Equations

We consider the solutions of the Nth order differential equations of the form:

$$F\left(x, y, \frac{dy}{dx}, \frac{d^2y}{dx^2}, \frac{d^3y}{dx^3}, \dots, \frac{d^{n-1}y}{dx^{n-1}}, \frac{d^n y}{dx^n}\right) = 0$$
(1-112)

Equation 1-112 has the highest derivative of the order n, and is ordinary because there is only one independent variable, x. We can obtain a unique solution when some additional information such as values of y(x) and its derivatives at some specific values of x are known. Therefore, for an Nth order equation, we require such N conditions to arrive at the unique solution y(x). If all N conditions are known at the same value of x, we can classify the problem as an initial value problem. However, when more than one value of x is involved, the problem is classified as a boundary value problem.

Solution of First Order Ordinary Differential Equations

We express a first order equation as:

$$\frac{\mathrm{d}y}{\mathrm{d}x} = f(x, y) \tag{1-113}$$

and we require a solution y(x) that satisfies Equation 1-113 and one initial condition. Here, we can subdivide the interval in the independent variable in x into steps over which a solution is required (a,b). The value of the exact solution y(x) is then approximated at N+1 evenly spaced values of x; i.e., $(x_0, x_1, \ldots, x_{n-1}, x_n)$. The step size, h, is expressed as:

and

 $x_i = x_0 + ih, \quad i = 0, 1, \dots n$

If we let the true solution y(x) be $y(x_i)$, and the computed approximations of y(x) at these same points be y_i , so that

$$\mathbf{y}_{i} = \mathbf{y}(\mathbf{x}_{i})$$

then, the exact derivative dy/dx can be approximated by $f(x_i, y_i)$ and represented as f_i such that

$$f_{i} = f(x_{i}, y_{i}) = f(x_{i}, y(x_{i}))$$
(1-114)

The difference between the computed value y_i and the true value $y(x_i)$ is ε_i and can be expressed as:

$$\varepsilon_i = y_i - y(x_i) \tag{1-115}$$

 ε_i is the local truncation error.

Taylor Series Expansion

We can develop a relation between x and y by finding the coefficients of the Taylor series. Expanding y about the point $x = x_0$, we obtain

$$y(x) = y(x_0) + f'(x_0)(x - x_0) + \frac{f''(x_0)(x - x_0)^2}{2!} + \frac{f'''(x_0)(x - x_0)^3}{3!} + \dots$$
(1-116)

where f'(x) represents $\frac{d}{dx}f(x)$

If we let $\mathbf{x} - \mathbf{x}_0 = \mathbf{h}$

We can express the series as:

$$y(x) = y(x_0) + hf'(x_0) + \frac{f''(x_0)h^2}{2!} + \frac{f'''(x_0)h^3}{3!} + \dots$$
(1-117)

Since $y(x_0)$ is our initial condition, the first term is known from the initial condition y(0) = 1. The error term of the Taylor series after the h⁴ term is

$$E = \frac{y^{v} \varepsilon h^{5}}{5!} \quad 0 < \varepsilon < h$$

Euler and Modified Euler Methods

Using the Taylor's series,

$$y(x_0 + h) = y(x_0) + hf'(x_0) + \frac{f''(\varepsilon)h^2}{2}$$
 (1-118)

 $\mathbf{x}_0 < \mathbf{\epsilon} < \mathbf{x}_0 + \mathbf{h}$

The value of $y(x_0)$ is given by the initial condition and $f'(x_0)$ is evaluated from $f(x_0,y_0)$, given by the differential equation

$$\frac{\mathrm{d}y}{\mathrm{d}x} = \mathrm{f}(x,y)$$

The Euler method can be expressed as:

$$y_{n+1} = y_n + hf'_n + O(h^2)$$
 error (1-119)

For the modified Euler method, we expand the Taylor series as

$$y_{n+1} = y_n + f'_n h + \frac{f''_n}{2} h^2 + \frac{f'''_n}{6} h^3, \quad x_n < \varepsilon < x_{n+h}$$
 (1-120)

Replacing the second derivative by the forward difference approximation for f" that is

$$f'' = \frac{f'_{n+1} - f'_n}{h}$$

having an error of O(h), we have

$$y_{n+1} = y_n + h \left\{ f'_n + \frac{1}{2} \left[\frac{f'_{n+1} - f'_n}{h} + O(h) \right] h \right\} + O(h^3)$$
(1-121)
$$y_{n+1} = y_n + h \left\{ f'_n + \frac{1}{2} f'_{n+1} - \frac{1}{2} f'_n \right\} + O(h^3)$$
$$= y_n + \frac{h(f'_n + f'_{n+1})}{2} + O(h^3)$$
(1-122)

Runge-Kutta Methods

The solution of a differential equation by direct Taylor's expansion cannot be easily obtained, if we retain the derivatives of higher order. We can develop one-step procedures that involve only first-order derivative evaluations and produce results equivalent in accuracy to higher order Taylor formulas. These algorithms are named the Runge-Kutta methods after the German mathematicians Runge and Kutta. The second-order Runge-Kutta algorithm for the first-order differential equation can be expressed as:

$$\frac{dy}{dx} = f(x, y)$$

$$k_{1} = hf(x_{i}, y_{i})$$

$$k_{2} = hf(x_{i} + h, y_{i} + k_{i})$$

$$y_{n+1} = y_{n} + \frac{1}{2}(k_{1} + k_{2}) + O(h^{3})$$
(1-123)

The third-order Runge-Kutta method is

$$k_{1} = hf(x_{i}, y_{i})$$

$$k_{2} = hf\left(x_{i} + \frac{h}{2}, y_{i} + \frac{k_{1}}{2}\right)$$

$$k_{3} = hf(x_{i} + h, y_{i} + 2k_{2} - k_{1})$$

$$y_{n+1} = y_{n} + \frac{1}{6}(k_{1} + 4k_{2} + k_{3}) + O(h^{4})$$
(1-124)

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The fourth-order Runge-Kutta method is widely used in computer solutions of differential equations.

$$k_{1} = hf(x_{i}, y_{i})$$

$$k_{2} = hf\left(x_{i} + \frac{h}{2}, y_{i} + \frac{k_{1}}{2}\right)$$

$$k_{3} = hf\left(x_{i} + \frac{h}{2}, y_{i} + \frac{k_{2}}{2}\right)$$

$$k_{4} = hf(x_{i} + h, y_{i} + k_{3})$$

$$y_{n+1} = y_{n} + \frac{1}{6}(k_{1} + 2k_{2} + 2k_{3} + k_{4}) + O(h^{5})$$
(1-125)

 k_1, k_2, k_3, k_4 are approximate derivative values computed on the interval $x_i \le x \le x_{i+1}$ and h is the step size.

$$\mathbf{x}_{i+1} = \mathbf{x}_i + \mathbf{h}$$

The local error term for the fourth-order Runge-Kutta is O(h⁵).

Runge-Kutta-Gill Method

Runge-Kutta-Gill method is the most widely used single-step method for solving ordinary differential equations.

$$k_{1} = hf(x_{i}, y_{i})$$

$$k_{2} = hf\left(x_{i} + \frac{h}{2}, y_{i} + \frac{1}{2}k_{1}\right)$$

$$k_{3} = hf\left(x_{i} + \frac{h}{2}, y_{i} + \left[\frac{-1}{2} + \frac{1}{\sqrt{2}}\right]k_{1} + \left[1 - \frac{1}{\sqrt{2}}\right]k_{2}\right)$$

$$k_{4} = hf\left(x_{i} + h, y_{i} - \frac{1}{\sqrt{2}}k_{2} + \left[1 + \frac{1}{\sqrt{2}}\right]k_{3}\right)$$

$$y_{n+1} = y_{n} + \frac{1}{6}\left(k_{1} + 2\left[1 - \frac{1}{\sqrt{2}}\right]k_{2} + 2\left[1 + \frac{1}{\sqrt{2}}\right]k_{3} + k_{4}\right) + O(h^{5})$$
(1-126)

Runge-Kutta-Gill method provides an efficient algorithm for solving a system of first-order differential equations and makes use of much less computer memory when compared with other numerical methods.

Runge-Kutta-Merson Method

Runge-Kutta-Merson outlines a process for deciding the step size for a better predetermined accuracy. For this method, five functions are evaluated at every step. The algorithm is

$$k_{1} = hf(x_{i}, y_{i})$$

$$k_{2} = hf\left(x_{i} + \frac{h}{3}, y_{i} + \frac{k_{1}}{3}\right)$$

$$k_{3} = hf\left(x_{i} + \frac{h}{3}, y_{i} + \frac{k_{1}}{6} + \frac{k_{2}}{6}\right)$$

$$k_{4} = hf\left(x_{i} + \frac{h}{2}, y_{i} + \frac{k_{1}}{8} + \frac{3k_{3}}{8}\right)$$

$$k_{5} = hf\left(x_{i} + h, y_{i} + \frac{k_{1}}{2} - \frac{3k_{3}}{2} + 2k_{4}\right)$$

$$y_{n+1} = y_{n} + \frac{1}{6}(k_{1} + 4k_{4} + k_{5}) + O(h^{5})$$
(1-127)

We can estimate the local error from a weighted sum of the individual estimate.

$$E = \frac{1}{30} (2k_1 - 9k_3 + 8k_4 - k_5)$$
(1-128)

Multistep Methods

The multistep methods use past values of y and y' = f(x,y) to construct a polynomial that approximates the derivative function and extrapolates this into the next interval. We can achieve an accurate estimate of y_{n+1} based on a polynomial that fits past values of the gradient. Also, we end with a slope equal to that at the predicted point. This forms the basis of predictor-corrector formulae.

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Fourth Order Milne's Method

Values of y_n , y_{n-1} , y_{n-2} and y_{n-3} are required to calculate y_{n+1} . Milne's method uses Newton-Cotes formula for the predictor and Simpson's rule for the corrector.

Predictor

$$y_{n+1}^{p} = y_{n-3} + \frac{4h}{3} [2f(x_{n}, y_{n}) - f(x_{n-1}, y_{n-1}) + 2f(x_{n-2}, y_{n-2})] + O(h^{5})$$
(1-129)

Corrector

$$y_{n+1} = y_{n-1} + \frac{h}{3} [f(x_{n+1}, y_{n+1}^{p}) + 4f(x_{n}, y_{n}) + f(x_{n-1}, y_{n-1})] + O(h^{5})$$
(1-130)

Local truncation error

$$O(h^5) = -\frac{h^5}{90} f^{v}(\epsilon), \quad x_{n-1} < \epsilon < x_{n+1}$$

Adams-Moulton Fourth-Step Method

Predictor

$$y_{n+1}^{p} = y_{n} + \frac{h}{24} \left[55f(x_{n}, y_{n}) - 59f(x_{n-1}, y_{n-1}) + 37f(x_{n-2}, y_{n-2}) - 9f(x_{n-3}, y_{n-3}) \right] + O(h^{5})$$
(1-131)
n = 3, 4, ... m - 1

Corrector

$$y_{n+1} = y_n + \frac{h}{24} \Big[9f(x_{n+1}, y_{n+1}^p) + 19f(x_n, y_n) - 5f(x_{n-1}, y_{n-1}) + f(x_{n-2}, y_{n-2}) \Big] + O(h^5) \quad n = 2, 3, \dots m - 1 \quad (1-132)$$

Local truncation error

$$O(h^5) = -\frac{19}{720} h^5 f^{v}(\epsilon), \quad x_{n-2} < \epsilon < x_{n+1}$$

The predictor calls for four previous values in Adams-Moulton and Milne's algorithms. We obtain these by the fourth-order Runge-Kutta method. Also, we can reduce the step size to improve the accuracy of these methods. Milne's method is unstable in certain cases because the errors do not approach zero as we reduce the step size, h. Because of this instability, the method of Adams-Moulton is more widely used.

The primary advantage of the single-step methods is that they are self starting. We can also vary the step sizes. In contrast, the multistep methods require a single-step formula to start the calculations. Step size variation is difficult. However, the efficiency of both the Milne's and Adams-Moulton methods is about twice that of the single-steps methods. We need two function evaluations per step in the former while four or five are required with the single step.

PROBLEMS AND SOLUTIONS

Problem 1-1

An experimental result in a liquid mixing experiment for the power correlation using a pitched-blade turbine shows that in the viscous regime, the power number is related to the Reynolds number by the following data:

Reynolds number, N _{Re}	Power number, Po	
$\frac{D^2 N \rho}{\mu}$	$\frac{Pg_{c}}{\rho N^{3}D^{5}}$	
1.0	50.0	
3.0	18.0	
5.0	11.0	
7.0	7.9	
9.0	6.2	
13.0	5.0	

Table P1-1

Use the regression analysis to determine the best fit for the data.

Solution

Program PROG11.FOR determines the slope, intercept, and correlation coefficient for the above experimental data. The results show that Equation Y=A+B*1/X gives the best fit with the slope B = 49.01 and a correlation coefficient r = 0.9998. Figure 1-7 gives a plot of Reynolds number against power number. The figure shows that Equation 1-3 gives a perfect fit of the experimental data.

Problem 1-2

In a fluid flow experiment, the volumetric rate of fluid through a pipe is dependent on the pipe diameter and slope by the following equation.

$$Q = C_0 D^{C_1} D^{C_2}$$
(1-133)

where Q =flow rate, ft^3/sec D =pipe diameter, ft

S = slope, ft/ft

Determine the flow rate of fluid for a pipe with a diameter of 3.25 ft and a slope of 0.025 ft.

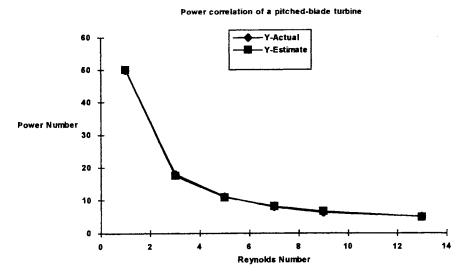


Figure 1-7. A characteristic impeller curve of power number as a function of Reynolds number.

Diameter ft X ₁	Slope, ft/ft X ₂	Flowrate, ft^3/s Y
1.0	0.001	1.5
2.5	0.005	9.0
3.0	0.010	25.0
4.0	0.010	5.0
1.0	0.050	30.0
3.5	0.050	100.0

Table P1-2

Program PROG12 determines the values of the coefficients C_0 , C_1 , and C_2 and the correlation coefficient. The coefficients are:

 $C_0 = 563.24$ $C_1 = 0.2725$ $C_2 = 0.8696$

The correlation coefficient r = 0.9045.

The predicted flow for a pipe with a diameter of 3.25 ft and a slope of 0.03 ft/ft is 36.81 ft^3/sec.

Problem 1-3

The following data are obtained from $y(x) = x^4 + 3x^3 + 2x^2 + x + 5$. Show that a fourth-degree polynomial provides the best least squares approximation to the given data. Determine this polynomial.

X	0.1	0.2	0.3	0.4	0.5	0.6	0.7
Y	5.123	5.306	5.569	5.938	6.437	7.098	7.949

Table P1-3

X	0.8	0.9
Υ	9.025	10.363

Program PROG13 calculates (i) the coefficients for each degree of the polynomial, (ii) the variance, (iii) error sum of squares, (iv) total sum of squares, (v) coefficient of determination, and (vi) the correlation coefficient. The program shows that the fourth degree gives the lowest value of the variance, and therefore shows the best fit. The results are:

(i) the variance	= 0.5557E-06
(ii) the error sum of squares	= 0.2223E-05
(iii) the total sum of squares	= 0.2620E+02
(iv) the correlation coefficient	= 1.00

The calculated polynomial is:

 $y(x) = 4.999 + 1.011x + 1.965x^2 + 3.047x^3 + 0.977x^4$

Problem 1-4

The final product from a chemical factory is made by blending four liquids (α , β , γ , δ) together. Each of these liquids contains four components A, B, C, and D. The product leaving the factory has to have a closely specified composition. Determine the relative quantities of α , β , γ , and δ required to meet the blend specifications in the following data:

		w/w c	omposition c		w/w composition of
Component	α	β	γ	δ	specification
Α	51.30	43.20	56.40	47.40	48.80
В	11.30	11.50	15.50	8.50	11.56
С	29.40	31.50	22.50	30.40	29.43
D	8.00	10.30	5.60	13.70	10.21

Table P1-4

Source: B.Sc. Final year 1978, Aston University, Birmingham, U.K.

Program PROG14 uses the Gaussian elimination method to determine the quantities of each of α , β , γ , and δ and required to meet the blending specification. Answers:

 $\alpha = 0.1172$ $\beta = 0.3789$ $\gamma = 0.2117$ $\delta = 0.3054$

Problem 1-5

A chemical reaction takes place in a series of four continuous stirred tank reactors arranged as shown. The chemical reaction is a first-order irreversible reaction of the type.

 $A \xrightarrow{k_1} B$

The conditions of temperature in each reactor are such that the values of k_i and V_i are given in Table P1-5. Figure 1-8 shows four continuous stirred tanks with recycle streams.

The following assumptions are:

- 1. The system is at steady state.
- 2. The reactions are in the liquid phase.

Reactor	Volume, V _i L	Rate Constant, k _i h−1
1	1000	0.1
2	1500	0.2
3	100	0.4
4	500	0.3

Table P1-5

Source: A. Constantinides [1], Applied Numerical Methods With Personal Computers, Courtesy of McGraw-Hill Book Co., 1987.

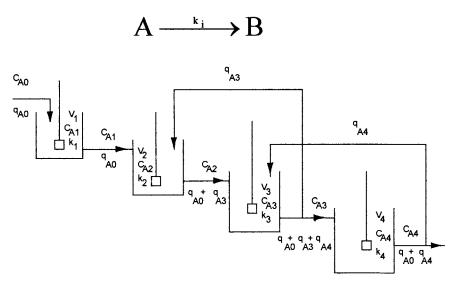


Figure 1-8. Chemical reaction with recycles in four continuous stirred tanks.

- 3. There is no change in volume or density of the liquid.
- 4. The rate of disappearance of component A in each in reactor is given by:

$$(-\mathbf{r}_{A}) = \mathbf{k}\mathbf{C}_{A}$$

Set up the material balance equations for each of the four reactors, and use the Gauss-Seidel method to determine the exit concentration from each reactor.

Solution

The general unsteady state material balance for each reactor is:

Input = output + disappearance + accumulation by reaction

Mass balance for reactor 1.

$$q_{A0}C_{A0} = q_{A0}C_{A1} + (-r_A)V_1 + V_1 \frac{dC_{A1}}{dt}$$

Because the system is at steady state, the accumulation is zero, the above equation becomes

$$q_{A0}C_{A0} = q_{A0}C_{A1} + k_1C_{A1}V_1$$

Mass balance for reactor 2.

$$q_{A0}C_{A1} + q_{A3}C_{A3} = (q_{A0} + q_{A3})C_{A2} + k_2C_{A2}V_2$$

Mass balance for reactor 3.

 $(q_{A0} + q_{A3})C_{A2} + q_{A4}C_{A4} = (q_{A0} + q_{A3} + q_{A4})C_{A3} + k_3V_3C_{A3}$

Mass balance for reactor 4.

$$(q_{A0} + q_{A4})C_{A3} = (q_{A0} + q_{A4})C_{A4} + k_4V_4C_{A4}$$

where

$$q_{A0} = 1000 \frac{\text{liters}}{\text{h}}, C_{A0} = 1 \frac{\text{mol}}{\text{liter}}$$

$$q_{A3} = 100 \frac{\text{liters}}{\text{h}}$$

$$q_{A4} = 100 \frac{\text{liters}}{\text{h}}$$

$$(1000)(1) = 1000C_{A1} + (0.1)(C_{A1})(1000)$$

$$1000C_{A1} + 100C_{A3} = 1100C_{A2} + (0.2)(C_{A2})(1500)$$

$$1100C_{A2} + 100C_{A4} = 1200C_{A3} + (0.4)(C_{A3})(100)$$

$$1100C_{A3} = 1100C_{A4} + (0.3)(C_{A4})(500)$$

Rearranging the above equations:

$$1100C_{A1} = 1000$$

$$1000C_{A1} -1400C_{A2} +100C_{A3} = 0$$

$$1100C_{A2} -1240C_{A3} +100C_{A4} = 0$$

$$1100C_{A3} -1250C_{A4} = 0$$

The above is a set of four simultaneous linear algebraic equations. This is a diagonal system of equations because the coefficients on the diagonal are larger in above value than the sum of the absolute values of the other coefficients. Therefore, the best method of solution for a diagonal system of linear algebraic equations is the Gauss-Seidel method. PROG15 is the Gauss-Seidel computer program for solving the above

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equations. The four material balance equations are rearranged to solve for the unknown on the diagonal position of each equation.

$$C_{A1} = \frac{1000}{1100}$$

$$C_{A2} = \frac{1000C_{A1} + 100C_{A3}}{1400}$$

$$C_{A3} = \frac{1100C_{A2} + 100C_{A4}}{1240}$$

$$C_{A4} = \frac{1100C_{A3}}{1250}$$

An initial guess of C_{A1} , C_{A2} , C_{A3} and $C_{A4} = 0.5$ is used to start the Gauss-Seidel iterative method. The method converges after five iterations to the solutions:

$$C_{A1} = 0.9091$$

 $C_{A2} = 0.6969$
 $C_{A3} = 0.6654$
 $C_{A4} = 0.5856$

Problem 1-6

The Colebrook implicit equation for the Darcy friction factor, f_D , for turbulent flow is:

$$F(f_{\rm D}) = \frac{1}{f_{\rm D}^{0.5}} + 0.86858 \ln \left[\frac{\epsilon / D}{3.7} + \frac{2.51}{N_{\rm Re} f_{\rm D}^{0.5}}\right]$$

Determine the friction factor for $N_{Re} = 184000$, $\varepsilon = 0.00015$ ft and D = 0.17225ft (2.067 inch).

Solution

PROGRAM PROG16 solves the above implicit equation. The program uses Newton-Raphson method to give $f_D = 0.02063$ after three iterations.

Problem 1-7

In a fire tube boiler where gas flows inside the tubes and a steam-water mixture flows on the outside, the heat transfer coefficient inside $h_i(Btu/ft^2hr.^{\circ}F)$ can be expressed as:

$$h_{i} = 2.44 \frac{w^{0.8} F_{i}}{d^{0.8}}$$

where factors F_1 , F_2 , and F_3 are:

$$F_{1} = \left(\frac{C_{p}}{\mu}\right)^{0.4} k^{0.6}$$
$$F_{2} = \left(\frac{C_{p}}{\mu}\right)^{0.3} k^{0.7}$$
$$F_{3} = \left(\frac{F_{2}}{C_{p}}\right) \mu^{=0.15}$$

Table P1-6 shows F_1 , F_2 , and F_3 for flue gas at a temperature range 200°F $\leq T \leq 1200$ °F. Determine the values of F_1 , F_2 , and F_3 if the film temperature is 575°F.

Temperature °F	F ₁	F ₂	F ₃
200	0.1700	0.0954	0.5851
300	0.1770	0.1015	0.6059
400	0.1835	0.1071	0.6208
600	0.1943	0.1170	0.6457
800	0.2051	0.1264	0.6632
1000	0.2136	0.1340	0.6735
1200	0.2216	0.1413	0.6849

Table P1-6

Source: V. Ganapathy, "Simplified Approach to Designing Heat Transfer Equipment," Courtesy of Chem. Eng., April 13, 1987, pp. 88–87.

PROG17 uses the interpolation methods of determine the values of F_1 , F_2 , and F_3 at t = 575°F. The values from the Stirling's central difference formula are:

 $F_1 = 0.1932$ $F_2 = 0.1160$ $F_3 = 0.6431$

These values give the best interpolation at $t = 575^{\circ}F$ rather than the Newton-Gregory's forward or backward interpolation formulae.

Problem 1-8

A tracer experiment was carried out in a nozzle type reactor of volume V = 5.13L with liquid rate at 2.9 l/min. Table P1-7 shows data for the exit age distribution $E(\theta)$ against the dimensionless residence time θ . Determine the area under the distribution curve.

θ	E(θ)	θ	Ε(θ)
0.000	0.000	1.243	0.403
0.113	0.308	1.356	0.355
0.226	0.995	1.469	0.313
0.339	0.876	1.582	0.275
0.452	0.786	1.695	0.237
0.565	0.720	1.808	0.213
0.678	0.663	1.921	0.171
0.791	0.606	2.034	0.142
0.904	0.545	2.147	0.123
1.017	0.497	2.260	0.109
1.130	0.450	2.373	0.095

Table P1-7

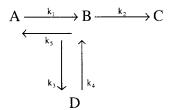
Source: A. K. Coker, Ph.D., "A Study of Fast Reactions in Nozzle Type Reactors," 1985, Aston University, Birmingham, U.K.

Program PROG18 uses Simpson's rule to compute the area of the exit age distribution of a tracer from a nozzle reactor.

The area under the distribution curve $\int_{0.0}^{2.373} E(\theta) = 1.0037$

Problem 1-9

Design of a batch reactor for complex reactions of the type



Determine the concentrations of A, B, C, and D over a period of ten minutes. The following rate constants and initial amounts for first-order reactions are:

Rate Constant	Concentration at time t = 0
	$\frac{\text{mol}}{\text{m}^3}$
k ₁ = 0.45	$C_{A0} = 9.90$
$k_2 = 0.16$	C _{B0} = 0.0
$k_3 = 0.12$	$C_{c0} = 0.0$
$k_4 = 0.08$	C _{D0} = 0.5
$k_5 = 0.10$	

Table P1-8

Solution

The mass balances for the batch reactor involving components A, B, C, and D are:

$$(-r_{A})_{net} = k_{1}C_{A} - k_{5}C_{B} = -\frac{dC_{A}}{dt}$$

$$(-r_{\rm B})_{\rm net} = (k_2 + k_3 + k_5)C_{\rm B} - k_1C_{\rm A} - k_4C_{\rm D} = -\frac{dC_{\rm B}}{dt}$$
$$(r_{\rm C})_{\rm net} = k_2C_{\rm B} = \frac{dC_{\rm C}}{dt}$$
$$(-r_{\rm D})_{\rm net} = k_4C_{\rm D} - k_3C_{\rm B} = -\frac{dC_{\rm D}}{dt}$$

Rearranging the above equations

$$\frac{dC_{A}}{dt} = k_{5}C_{B} - k_{1}C_{A}$$
$$\frac{dC_{B}}{dt} = k_{1}C_{A} + k_{4}C_{D} - (k_{2} + k_{3} + k_{5})C_{B}$$

Table	P1	-9
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Runge-Kutta Fourth Order Method for a System of Ordinary Differential Equations

TIME	CONC. CA	CONC. CB	CONC. CC	CONC. CD
.00	9.90	.00	.00	.50
.50	8.15	1.65	.07	.54
1.00	6.77	2.72	.25	.66
1.50	5.68	3.39	.50	.83
2.00	4.82	3.77	.78	1.03
2.50	4.12	3.95	1.09	1.23
3.00	3.55	4.00	1.41	1.44
3.50	3.09	3.94	1.73	1.64
4.00	2.71	3.83	2.04	1.82
4.50	2.39	3.68	2.34	1.99
5.00	2.12	3.51	2.63	2.14
5.50	1.89	3.33	2.90	2.27
6.00	1.69	3.15	3.16	2.39
6.50	1.52	2.97	3.41	2.49
7.00	1.38	2.80	3.64	2.58
7.50	1.25	2.64	3.86	2.65
8.00	1.14	2.49	4.06	2.71
8.50	1.04	2.34	4.26	2.76
9.00	.96	2.21	4.44	2.79
9.50	.88	2.09	2.61	2.82
10.00	.81	1.97	4.77	2.84

$$\frac{dC_{\rm C}}{dt} = k_2 C_{\rm B}$$
$$\frac{dC_{\rm D}}{dt} = k_3 C_{\rm B} - k_4 C_{\rm D}$$

where $F(1) = dC_A/dt$, $F(2) = dC_B/dt$, $F(3) = dC_C/dt$ and $F(4) = dC_D/dt$ and $X(1) = C_A$, $X(2) = C_B$, $X(3) = C_C$ and $X(4) = C_D$.

Thus, we can express the differential equations in the form of x-arrays as:

$$\begin{split} F(1) &= k_5 \cdot X(2) - k_1 \cdot X(1) \\ F(2) &= k_1 \cdot X(1) + k_4 \cdot X(4) - (k_2 + k_3 + k_5) \cdot X(2) \\ F(3) &= k_2 \cdot X(2) \\ F(4) &= k_3 \cdot X(2) - k_4 \cdot X(4) \end{split}$$

Program PROG19 uses the Runge-Kutta fourth-order to solve the differential equations F(1), F(2), F(3), and F(4). Table P1-9 gives the computer printouts of C_A , C_B , C_C , and C_D from t= 0 to t=10 minutes. Figure 1-9 shows the profiles of concentrations from the start of the batch reaction to the final time of ten minutes.

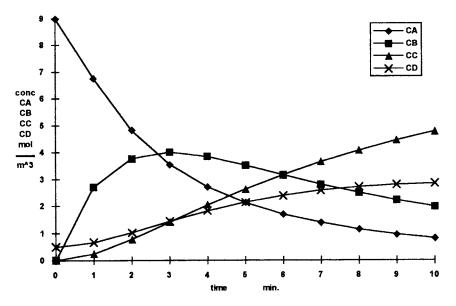


Figure 1-9. Concentration profiles of A, B, C, and D as a function of time in a batch reactor.

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PROGRAM PROG11

*********** С * THE PROGRAM DETERMINES A RELATIONSHIP BETWEEN TWO VARIABLES С i.e. ONE DEPENDENT AND ONE INDEPENDENT VARIABLE. THE PROGRAM * * 00000 * SELECTS THE CURVE OF BEST FIT BY TESTING THE DATA AGAINST × EQUATIONS OF THE FORM: Y=a+b.F(x) + THE PROGRAM GIVES THE CONSTANTS FOR THE EQUATION AND THE * * • CORRELATION COEFFICIENT. С THIS PROGRAM WILL CORRELATE X-Y DATA FOR THE FOLLOWING EQUATIONS: с С 1. Y=a+b.X С 2. Y=a+b.X^2 с 3. Y=a+b.X^0.5 c c Y=a.exp(b.X)4. 5. Y=a+b.ln(X)С 6. Y=a.X^b С Y=a+b.(1/X) 7. с 8. 1/Y = a + b.(1/X)С VECTOR OF INDEPENDENT VARIABLE х = VECTOR OF DEPENDENT VARIABLE С Y = c c YCAL = VECTOR OF THE ESTIMATED DEPENDENT VARIABLE NUMBER OF DATA N = A, B = c CONSTANTS FOR THE EQUATION с COR = CORRELATION COEFFICIENT с DIMENSION X(1:50), Y(1:50) DIMENSION YCAL1(1:50), YCAL2(1:50), YCAL3(1:50), YCAL4(1:50) DIMENSION YCAL5(1:50), YCAL6(1:50), YCAL7(1:50), YCAL8(1:50) COMMON/DATA/N,X,Y COMMON/DATA1/YCAL1,A1,B1 COMMON/DATA2/YCAL2,A2,B2 COMMON/DATA3/YCAL3, A3, B3 COMMON/DATA4/YCAL4, A4, B4 COMMON/DATA5/YCAL5, A5, B5 COMMON/DATA6/YCAL6, A6, B6 COMMON/DATA7/YCAL7, A7, B7 COMMON/DATA8/YCAL8, A8, B8 INTEGER N OPEN (UNIT=3, FILE='DATA11.DAT', STATUS='OLD', ERR=18) OPEN(UNIT=1, FILE='PRN') READ (3, *, ERR=19)N
READ (3, *, ERR=19)(X(I), Y(I), I=1,N)

GO TO 2

18 111 19 222	WRITE (1, 111) FORMAT (//,6X, 'FILE DOES NOT EXIST') GO TO 999 WRITE (1, 222) FORMAT (//,6X, 'ERROR MESSAGE IN THE DATA VALUE') GO TO 999
2 100 *	<pre>WRITE(1, 100) FORMAT(//,1H0,25X,'CURVE FITTING FOR TWO VARIABLES',</pre>
110	WRITE(1, 110) FORMAT(//,1H0,25X,'X',15X,'Y',/1H ,78(1H-))
	DO 10 I=1,N
120 10	WRITE(1, 120)I,X(I),Y(I) FORMAT(1H0,4X,I4,6X,F14.3,6X,E14.6) CONTINUE
125	WRITE (1, 125) FORMAT (1H ,78(1H-))
130	WRITE(1, 130) FORMAT(//, 1H ,25X,'THE RESULTS ARE:',/1H ,78(1H*))
140 *	<pre>WRITE(1, 140) FORMAT(/,10X,'EQUATION',10X,'REGRESSION COEFFICIENTS', 5X,'CORRELATION')</pre>
150 *	<pre>WRITE(1, 150) FORMAT (28X,'INTECEPT: A',5X,'SLOPE: B',4X,'COEFFICIENT',</pre>
	J=1
4	GO TO (5,15,25,35,45,55,65,75),J
с	CURVE FITTING FOR : Y=A+BX
5	CALL LINFIT(X, Y, N, A1, B1, COR1)
160	WRITE (1, 160)A1, B1, COR1 FORMAT(2X,'1.',5X,'Y=A+BX',10X,3(E10.4,5X))
30	DO 30 I=1,N YCAL1(I)=A1+B1*X(I) CONTINUE
	J=J+1
	IF (J.EQ. 2) THEN GO TO 4 ELSE

GO TO 800 ENDIF CURVE FITTING FOR Y=A+BX^2 С DO 40 I=1,N X(I)=X(I)*X(I) 15 40 CONTINUE CALL LINFIT(X, Y, N, A2, B2, COR2) WRITE(1, 170) A2, B2, COR2 FORMAT (2X,'2.',5X,'Y=A+B*X^2',7X,3(E10.4,5X)) 170 DO 44 I=1,N YCAL2(I)=A2+B2*X(I)**2 CONTINUE 44 J=J+1 IF (J .EQ. 3) THEN GO TO 4 ELSE GO TO 800 ENDIF с CURVE FITTING FOR Y=A+B.X^0.5 DO 50 I=1,N X(I)=X(I)**0.5 25 CONTINUE 50 CALL LINFIT(X, Y, N, A3, B3, COR3) WRITE(1, 180) A3, B3, COR3 FORMAT(2X,'3.',5X,'Y=A+B*X^0.5',5X,3(E10.4,5X)) 180 DO 54 I=1,N YCAL3(I)=A3+B3*X(I)**0.5 54 CONTINUÉ J=J+1 IF (J.EQ. 4) THEN GO TO 4 ELSE GO TO 800 ENDIF С CURVE FITTING FOR Y=A.EXP(B.X) с LINEARIZE TO Ln(Y)=Ln(A)+BX

35 DO 60 I=1,N

```
Y(I) = ALOG(Y(I))
 60
         CONTINUE
         CALL LINFIT(X, Y, N, A4, B4, COR4)
         A4=EXP(A4)
         WRITE(1, 190) A4, B4, COR4
FORMAT(2X,'4.',5X,'Y=A*EXP(B*X)',4X,3(E10.4,5X))
 190
         DO 61 I=1,N
         Y(I) = EXP(\dot{Y}(I))
         YCAL4(I)=A4*EXP(B4*X(I))
         CONTINUE
 61
         J=J+1
         IF (J .EQ. 5) THEN
GO TO 4
              ELSE
              GO TO 800
         ENDIF
С
         CURVE FITTING FOR Y=A+B.Ln(X)
 45
         DO 70 I=1,N
         X(I) = ALOG(X(I))
 70
         CONTINUE
         CALL LINFIT(X, Y, N, A5, B5, COR5)
         WRITE(1, 200)A5, B5, COR5
FORMAT(2X,'5.',5X,'Y=A+B*Ln(X)',5X,3(E10.4,5X))
 200
         DO 71 I=1,N
X(I)=EXP(X(I))
         YCAL5(I)=A5+B5*ALOG(X(I))
 71
         CONTINUE
         J=J+1
         IF (J.EQ. 6) THEN
GO TO 4
              ELSE
              GO TO 800
         ENDIF
         CURVE FITTING FOR Y=A.X^B
C
C
         LINEARIZE TO Ln(Y)=Ln(A)+B*Ln(X)
         DO 80 I=1,N
 55
         X(I) = ALOG(X(I))
         Y(I) = ALOG(Y(I))
 80
         CONTINUE
         CALL LINFIT(X, Y, N, A6, B6, COR6)
         A6=EXP(A6)
```

210	WRITE(1,210)A6, B6, COR6 FORMAT(2X,'6.',5X,'Y=A*X^B',9X,3(E10.4,5X))
85	DO 85 I=1,N X(I)=EXP(X(I)) Y(I)=EXP(Y(I)) YCAL6(I)=A6*(X(I)**B6) CONTINUE
	J=J+1 IF (J .EQ. 7) THEN GO TO 4 ELSE GO TO 800 ENDIF
с	CURVE FITTING FOR Y=A+B.1/X
65	DO 90 I=1,N
90	X(I)=1/X(I) CONTINUE CALL INFIT(Y, Y, N, A7, B7, COP7)
220	CALL LINFIT(X, Y, N, A7, B7, COR7) WRITE(1,220)A7, B7, COR7 FORMAT(2X,'7.',5X,'Y=A+B*1/X',7X,3(E10.4,5X))
95	DO 95 I=1,N X(I)=1/X(I) YCAL7(I)=A7+B7*1/X(I) CONTINUE
	J=J+1
	IF (J.EQ. 8) THEN GO TO 4 ELSE GO TO 800 ENDIF
с	CURVE FITTING FOR 1/Y=A+B.1/X
75	DO 96 I=1,N X(I)=1/X(I) Y(I)=1/Y(I)
96	CONTINUE
	CALL LINFIT(X, Y, N, A8, B8, COR8)
230	WRITE(1, 230)A8, B8, COR8 FORMAT(2X,'8.',5X,'1/Y=A+B*1/X',5X,3(E10.4,5X))
	DO 97 I=1,N X(I)=1/X(I) Y(I)=1/Y(I) YCAL8(I)=X(I)/(A8*X(I)+B8)

```
97
       CONTINUE
 800
       CALL COMP(X, Y, N, COR1, COR2, COR3, COR4, COR5, COR6, COR7, COR8)
       CLOSE (3, STATUS='KEEP')
       CLOSE(1)
 999
       STOP
       END
с
с
с
с
        THIS PROGRAM PERFORMS A LINEAR REGRESSION ANALYSIS ON A SET
        OF X-Y VALUES.
        *********
       SUBROUTINE LINFIT(X, Y, N, A,B,COR)
DIMENSION X(1:50),Y(1:50)
       INTEGER N
       SUMX=0.0
       SUMY=0.0
       SUMXY=0.0
       SUMX2=0.0
       SUMY2=0.0
       DO 10 I=1,N
       XI = X(I)
       YI=Y(I)
       SUMX=SUMX+XI
       SUMY=SUMY+YI
       SUMXY=SUMXY+XI*YI
       SUMX2=SUMX2+XI*XI
       SUMY2=SUMY2+YI*YI
 10
       CONTINUE
       SXX=N*SUMX2-SUMX*SUMX
       SXY=N*SUMXY-SUMX*SUMY
       SYY=N*SUMY2-SUMY*SUMY
       CALCULATE THE CONSTANTS FOR THE EQUATION
с
       B=SXY/SXX
       XMEAN=SUMX/N
       YMEAN=SUMY/N
с
       THIS PRINTS THE RESULTS ON THE SCREEN.
       WRITE(*, 50)XMEAN,YMEAN
FORMAT(6X,'XMEAN:',F12.3,3X,'YMEAN:',F12.4)
 50
       A=YMEAN-B*XMEAN
       CALCULATE THE CORRELATION COEFFICIENT
с
       COR=ABS(SXY/SQRT(SXX*SYY))
       WRITE(*, 60)A, B, COR
```

60 FORMAT(6X,'A=',F12.3,3X,'B=',F12.3,3X,'COR=',F12.4)

RETURN END

00000 THIS PROGRAM COMPARES THE VALUES OF THE CORRELATION COEFFICIENT OF THE REGRESSION ANALYSES SUBROUTINE COMP(X, Y, N, COR1,COR2,COR3,COR4,COR5,COR6,COR7,COR8) DIMENSION X(1:50), Y(1:50) DIMENSION YCAL1(1:50), YCAL2(1:50), YCAL3(1:50), YCAL4(1:50) DIMENSION YCAL5(1:50), YCAL6(1:50), YCAL7(1:50), YCAL8(1:50) INTEGER N COMMON/DATA1/YCAL1,A1,B1 COMMON/DATA2/YCAL2,A2,B2 . COMMON/DATA3/YCAL3,A3,B3 COMMON/DATA4/YCAL4,A4,B4 COMMON/DATA5/YCAL5,A5,B5 COMMON/DATA6/YCAL6, A6, B6 COMMON/DATA7/YCAL7, A7, B7 COMMON/DATA8/YCAL8,A8,B8 С FIND THE MAXIMUM VALUE OF THE CORRELATION COEFFICIENTS CORR=AMAX1(COR1,COR2,COR3,COR4,COR5,COR6,COR7, COR8) WRITE(1,100)CORR 100 FORMAT(//,2X,'MAXIMUM CORRELATION COEFFICIENT:',F12.4) COMPARE THIS VALUE WITH THE CALCULATED VALUES OF THE C C REGRESSION ANALYSES. IF (CORR .EQ. COR1) THEN CALL OUTPUT(X, Y, YCAL1, N, A1, B1, COR1) GO TO 10 ELSEIF (CORR .EO. COR2) THEN CALL OUTPUT(X, Y, YCAL2, N, A2, B2, COR2) GO TO 10 ELSEIF (CORR .EQ. COR3) THEN CALL OUTPUT(X, Y, YCAL3, N, A3, B3, COR3) GO TO 10 ELSEIF (CORR .EQ. COR4) THEN CALL OUTPUT (X, Y, YCAL4, N, A4, B4, COR4) GO TO 10 ELSEIF (CORR .EQ. COR5) THEN CALL OUTPUT(X, Y, YCAL5, N, A5, B5, COR5) GO TO 10 ELSEIF (CORR .EQ. COR6) THEN CALL OUTPUT(X, Y, YCAL6, N, A6, B6, COR6)

	GO TO 10
	ELSEIF (CORR .EQ. COR7) THEN CALL OUTPUT (X, Y, YCAL7, N, A7, B7, COR7) GO TO 10
	ELSE IF (CORR .EQ. COR8) THEN CALL OUTPUT(X, Y, YCAL8, N, A8, B8, COR8) ENDIF
10	RETURN END
С С С С С С С	**************************************
	SUBROUTINE OUTPUT (X, Y, YCAL, N, A, B, COR) DIMENSION X(1:50), Y(1:50), YCAL(1:50) INTEGER N
100 *	<pre>WRITE(1, 100) FORMAT(/,1H0, 21X,'X-ACTUAL', 13X,'Y-ACTUAL', 11X,'Y-ESTIMATE',</pre>
110 10	WRITE(1, 110) I, X(I), Y(I), YCAL(I) FORMAT(1H0,4X,I4,6X,F14.3,2(6X,F14.6)) CONTINUE
120	WRITE (1, 120) Format (1H ,78(1H-))
130 * *	<pre>WRITE (1, 130)A, B, COR FORMAT(//,1H0,6X,'CONSTANTS FOR THE EQUATION:',/1H0,6X,</pre>

DATA11.DAT				
6				
1.0	50.0			
3.0	18.0			
5.0	11.0			
7.0	7.90			
9.0	6.20			
13.0	5.0			

CURVE FITTING FOR TWO VARIABLES				
	х	У		
1	1.000	.500000E+02		
2	3.000	.180000E+02		
3	5.000	.110000E+02		
4	7.000	.790000E+01		
5	9,000	.620000E+01		
6	13.000	.500000E+01		

THE RESULTS ARE:

	EQUATION	INTECEPT:		CORRELATION COEFFICIENT	
1.	Y=A+BX	.3577E+02	3066E+01	.7735E+00	
2.	Y=A+B*X^2	.2553E+02	1650E+00	.6036E+00	
з.	Y=A+B*X^0.5	.3577E+02	3066E+01	.7735E+00	
4.	Y=A*EXP(B*X)	.3605E+02	1792E+00	.9150E+00	
5.	Y = A + B + Ln(X)	.4416E+02	1772E+02	.9496E+00	
6.	Y=A*X^B	.4913E+02	9203E+00	.9986E+00	
7.	Y=A+B*1/X	.1123E+01	.4901E+02	.9998E+00	
8.	1/Y=A+B*1/X	.1575E+00	1559E+00	.8137E+00	

MAXIMUM CORRELATION COEFFICIENT: .9998

	X-ACTUAL	Y-ACTUAL	Y-ESTIMATE
1	1.000	50.000010	50.130660
2	3.000	18.000000	17.459000

3	5.000	11.000000	10.924670	
4	7.000	7.900001	8.124240	
5	9.000	6.200000	6.568446	
6	13.000	5.000000	4.892976	

CONSTANTS FO	OR THE EQUATION:		
A:	1.1232		
в:	49.0075		
CORRELATION	COEFFICIENT:	.9998	

***** с с с с LINEAR MULTIPLE REGRESSION ANALYSIS THE PROGRAM PERFORMS A LINEAR MULTIPLE REGRESSION ANALYSIS INVOLVING TWO OR MORE INDEPENDENT VARIABLES AND ONE DEPENDENT с с с с VARIABLE USING THE GAUSS-ELIMINATION METHOD. THE LINEAR MULTIPLE REGRESSION EQUATION IS OF THE FORM Y=Co+C1*x1+C2*x2+....+Ck*Xk C C C C C C C C DEPENDING ON THE NUMBER OF THE INDEPENDENT VARIABLES, SIMULTANEOUS SOLUTION BY GAUSS ELIMINATION IS EMPLOYED TO CALCULATE THE UNKNOWN REGRESSION CO-EFFICIENTS. C C C C C C C C ******* X1,....Xk = THE INDEPENDENT VARIABLES = THE DEPENDENT VARIBALE Y 00000000 CO,C1,..Ck = THE UNKNOWN REGRESSION COEFFICIENTS k = THE NUMBER OF INDEPENDENT VARIABLES. = THE NUMBER OF DATA VALUES = THE NUMBER OF EQUATIONS N N1 č DIMENSION A(1:50,1:50), B(1:50), X1(1:50), X2(1:50) DIMENSION Y(1:50), YCAL(1:50) REAL MEANY DATA N/6/, N1/3/ DATA X1(1)/1.00/, X2(1)/0.001/, Y(1)/1.5/ DATA X1(2)/2.50/, X2(2)/0.005/, Y(2)/9.0/ DATA X1(3)/3.00/, X2(3)/0.010/, Y(3)/25.0/ DATA X1(4)/4.00/, X2(4)/0.010/, Y(4)/5.0/ DATA X1(5)/1.00/, X2(5)/0.050/, Y(5)/30.0/ DATA X1(6)/3.50/, X2(6)/0.050/, Y(6)/100.0/ THIS OPEN STATEMENT PRINTS THE RESULTS ONTO A PRINTER С OPEN (UNIT=1, FILE='PRN') LINEAR MULTIPLE REGRESSION ANALYSIS FOR AN EQUATION C C Y=C0+C1*X1+C2*X2 WRITE(1, 100) 100 FORMAT(//,20X,'LINEAR MULTIPLE REGRESSION ANALYSIS',/1H ,20X, *'FOR AN EQUATION : Y=CO+C1*X1+C2*X2',/1H ,74(1H*)) WRITE(1, 110) 110 FORMAT(/,1H0,25X,'X1',15X,'X2',15X,'Y',/,1H ,74(1H-)) DO 20 I=1,N WRITE(1, 115)I,X1(I),X2(I),Y(I) 115 FORMAT(6X, I4, 3(6X, F12.4)) 20 CONTINUE

```
WRITE (1, 120)
120 FORMAT (1H ,74(1H-))
C
       LINEARIZE BOTH THE INDEPENDENT AND DEPENDENT VARIABLES.
       DO 12 J = 1, N
       CONTINUE
 12
       WRITE(1,121)
      FORMAT(/,1H',20X,'LINEARIZED INDEPENDENT AND DEPENDENT VARIABLES')
WRITE(1,122)
 121
 122 FORMAT(/,3X,70(1H*)//,1H0,25X,'X1',15X,'X2',15X,'Y',/,1H ,74(1H-))
 DO 15 I = 1,N
WRITE(1, 125)I, X1(I), X2(I), Y(I)
125 FORMAT(6X,I4,3(6X,F12.4))
 15
       CONTINUE
 WRITE (1, 126)
126 FORMAT (1H ,74(1H-))
       WRITE(1, 130)N1
 130 FORMAT(//,1H0,6X,'THE NUMBER OF EQUATIONS:',3X,14)
с
      CALCULATE THE COEFFICIENTS OF MATRIX : A AND R.H.S. VECTOR : Y
   25 SX1=0.0
      SX2=0.0
      SX1X2=0.0
      SX1SQ=0.0
      SX2SQ=0.0
      SY=0.0
      SX1Y=0.0
      SX2Y=0.0
      DO 30 J=1,N
      SX1=SX1+X1(J)
      SX2=SX2+X2(J)
      SX1X2=SX1X2+X1(J)*X2(J)
      SX1SQ=SX1SQ+X1(J)**2
SX2SQ=SX2SQ+X2(J)**2
      SY=SY+Y(J)
      SX1Y=SX1Y+X1(J)*Y(J)
      SX2Y=SX2Y+X2(J)*Y(J)
   30 CONTINUE
      MEANY=SY/N
      WRITE(*, 140)SX1,SX2,SY
  140 FORMAT(3X, 'SUMX1=', F10.4, 6X, 'SUMX2=', F10.4, 6X, 'SUMY=', F10.4)
с
      AUGMENT THE COEFFICIENTS OF MATRIX A WITH THE R.H.S. VECTOR Y
```

A(1,1) = FLOAT(N)

```
A(2,1)=SX1
A(3,1)=SX2
       A(1,2) = A(2,1)
       A(2,2) = SX1SQ
       A(3,2)=SX1X2

A(1,3)=A(3,1)

A(2,3)=A(3,2)
       A(3,3)=SX2SQ
       A(1,4)=SY
       A(2,4) = SX1Y
       A(3,4)=SX2Y
       NUMBER OF EQUATION: N1
С
       M≈N1+1
       CALL GAUSS(A,B,N1,M,TEST)
       IF (TEST .EQ. 0.0) THEN
GO TO 40
       ELSE
  WRITE(1, 160)
160 FORMAT(//,1H0,6X,'ERROR FLAG')
       ENDIF
       GO TO 999
  40 WRITE(1, 165)
165 FORMAT(//,1H0,74(1H*))
       CO=B(1)
       C1 = B(2)
       C2=B(3)
       CALCULATE THE ESTIMATED VALUES OF THE DEPENDENT VARIABLES
С
       DO 45 I=1,N
       YCAL(I) = CO + C1 * X1(I) + C2 * X2(I)
    45 CONTINUE
       CALCULATE THE CORRELATION COEFFICIENT: COR
С
с
       REGRESSION SUM OF SQUARES: SSR
¢
       TOTAL SUM OF SQUARES: SST
с
       ERROR (OR RESIDUAL) SUM OF SQUARES: SSE
       SSR=0.0
       SST=0.0
       SSE=0.0
       DO 50 I=1,N
       SSR=SSR+(YCAL(I)-MEANY)**2
SST=SST+(Y(I)-MEANY)**2
SSE=SSE+(Y(I)-YCAL(I))**2
   50 CONTINUE
С
С
       THE CORRELATION COEFFICIENT: COR
       COR=SQRT(1.0-(SSE/SST))
       CALL OUTPUT(X1,X2,Y,YCAL,B,N,SSR,SSE,SST,COR)
       CLOSE (UNIT= 1)
 999
       STOP
       END
```

```
0000000
               *****************
                                                                     *****
      THIS PROGRAM SOLVES A SET OF LINEAR EQUATIONS USING THE
GAUSSIAN ELIMINATION METHOD. A SEARCH IS MADE IN EACH COLUMN
      FOR THE LARGEST ELEMENT.
       SUBROUTINE GAUSS(A,X,N,NP1,TEST)
      DIMENSION A(1:50,50),X(1:50)
      NM1=N-1
      DO 10 I=1,NM1
      KP1=I+1
      L=I
      DO 20 J=KP1,N
      IF(ABS(A(J,I)) . LE. ABS(A(L,I))) THEN
      GO TO 20
      ELSE
      L≖J
      ENDIF
   20 CONTINUE
      IF(L .EQ. I) THEN
GO TO 30
      ELSE
      IF(ABS(A(L,I)) .LE. 1E-7) THEN
      GO TO 40
      ELSE
      ENDIF
с
с
с
      INTERCHANGE THE VALUES
      DO 50 J=1,NP1
      TEMP=A(I,J)
      A(I,J)=A(L,J)
A(L,J)=TEMP
   50 CONTINUE
      ENDIF
CCCC
      ELIMINATE ALL ELEMENTS IN THE COLUMN BELOW MAIN DIAGONAL
   30 D=A(I,I)
DO 60 J=1,NP1
      A(I,J)=A(I,J)/D
   60 CONTINUE
      DO 70 L=KP1,N
      E=A(L,I)
      DO 80 J=1,NP1
      A(L,J)=A(L,J)-E*A(I,J)
   80 CONTINUE
   70 CONTINUE
   10 CONTINUE
      IF(ABS(A(N,N)) .LE. 1E-7) THEN
      GO TO 40
      ELSE
С
      BACK SUBSTITUTION
      X(N) = A(N, NP1) / A(N, N)
      ENDIF
```

```
DO 85 J=1,NM1
       J1=N-J
       X(J1)=A(J1,NP1)
       DO 90 K=J1+1,N
       X(J1) = X(J1) - \dot{A}(J1,K) * X(K)
   90 CONTINUE
   85 CONTINUE
       TEST=0.0
       RETURN
   40 TEST=1.0
       RETURN
       END
С
       С
       THIS PROGRAM GIVES THE RESULTS OF THE REGRESSION ANALYSIS WITH
с
       THE ESTIMATED VALUE OF THE DEPENDENT VARIABLE, THE CONSTANTS
ċ
       FOR THE EQUATION AND THE CORRELATION COEFFICIENT.
č
       SUBROUTINE OUTPUT(X1,X2,Y,YCAL,B,N,SSR,SSE,SST,COR)
       DIMENSION X1(1:50), X2(1:50), Y(1:50), YCAL(1:50), B(1:50)
       WRITE(1, 100)
  100 FORMAT(//,1H0,18X,'X1-ACTUAL',7X,'X2-ACTUAL',7X,'Y-ACTUAL',7X,
      *'Y-ESTIMATE',/,1H ,74(1H-))
       DO 10 I=1,N
       WRITE(1,110)I,X1(I),X2(I),Y(I),YCAL(I)
  110 FORMAT(1H0,4X,14,6X,F10.3,6X,F10.3,6X,F10.4,6X,F10.4)
   10 CONTINUE
       WRITE (1, 115)
 115 FORMAT (1H ,74(1H-))
       CO=EXP(B(1))
       C1=B(2)
       C2=B(3)
       CALCULATE THE PREDICTED FLOW FOR A PIPE WITH A DIAMETER OF 3.25ft.
С
       AND A SLOPE OF 0.03 ft./ft.
С
c
       Q=C0.D^C1.S^C2
       Q=CO*(3.25**C1)*(0.03**C2)
       WRITE(1, 120)CO,C1,C2
  wRITE(1, 120)CO,C1,C2
120 FORMAT(//,1H0,6X,'COEFFICIENTS FOR THE EQUATION:',/1H0,6X,
*'CO=',3X,F12.4,/1H0,6X,'C1=',3X,F12.4,/1H0,6X,'C2=',3X,F12.4)
wRITE(1,130)SSR,SSE,SST,COR
130 FORMAT(/,1H0,6X,'REGRESSION SUM OF SQUARES:',F12.4,/1H0,6X,
'WRITE(1,120)CO,CONTRACT,'SOUND OF SQUARES:',F12.4,/1H0,6X,
     *'ERROR SUM OF SQUARES:',F12.4,/1H0,6X,
*'TOTAL SUM OF SQUARES:',F12.4,/1H0,6X,
      *'CORRELATION COEFFICIENT:', F12.4)
 WRITE(1, 140)Q
140 FORMAT(//,1H ,6X,'THE PREDICTED FLOW FOR A PIPE WITH A DIAMETER',
 */1H ,6X,'OF 3.25ft AND SLOPE OF 0.03 ft/ft IS:',F8.2,1X,'ft.^3/s')
       WRITE (1, 150)
 150 FORMAT (1H ,74(1H-))
       RETURN
       END
```

LINEAR MULTIPLE REGRESSION ANALYSIS FOR AN EQUATION : Y=CO+C1*X1+C2*X2

	Xl	X2	Y	
1	1.0000	.0010	1.5000	
2	2.5000	.0050	9.0000	
3	3.0000	.0100	25.0000	
4	4.0000	.0100	5.0000	
5	1.0000	.0500	30.0000	
6	3.5000	.0500	100.0000	
******	LINEARIZED IN	IDEPENDENT AND DEPI	ENDENT VARIABLES	***
*****	LINEARIZED IN	IDEPENDENT AND DEPI	ENDENT VARIABLES	***
••••••••••••••••••••••••••••••••••••••	X1	X2	**************************************	***
**************************************	X1 .0000	X2 -6.9078	¥*************************************	***
2	X1 .0000 .9163	X2 	¥*************************************	***
2 3	X1 .0000 .9163 1.0986	X2 -6.9078 -5.2983 -4.6052	¥ ************************************	***
2	X1 .0000 .9163	X2 	¥*************************************	***

THE NUMBER OF EQUATIONS: 3

	X1-ACTUAL	X2-ACTUAL	Y-ACTUAL	Y-ESTIMATE
1	.000	-6.908	.4055	.3269
2	.916	-5.298	2.1972	1.9761
3	1.099	-4.605	3.2189	2.6286
4	1.386	-4.605	1.6094	2.7069
5	.000	-2.996	3.4012	3.7287
6	1.253	-2.996	4.6052	4.0701

COEFFICIENTS FOR THE EQUATION: CO= 563.2403 C1= .2725 C2= .8696 REGRESSION SUM OF SQUARES: 8.9989 ERROR SUM OF SQUARES: 2.0016 TOTAL SUM OF SQUARES: 11.0006 CORRELATION COEFFICIENT: .9045

THE PREDICTED FLOW FOR A PIPE WITH A DIAMETER OF 3.25ft AND SLOPE OF 0.03 ft/ft IS: 36.81 ft.^3/s

000000	**************************************
	PARAMETERS ARE:- X,Y = ARRAY OF X AND Y VALUES N = NUMBER OF DATA PAIRS MS, MF = THE RANGE OF DEGREE OF POLYNOMIALS TO BE COMPUTED A = AUGMENTED ARRAY OF THE COEFFICIENTS OF THE NORMAL = EQUATIONS C = ARRAY OF COEFFICIENTS OF THE LEAST SQUARES POLYNOMIAL
	DIMENSION X(1:100), Y(1:100), C(1:10), A(1:10,1:11) DIMENSION XN(1:100)
	OPEN (UNIT = 1, FILE='PRN')
	DATA N/9/ DATA X(1)/0.1/, Y(1)/5.123/, X(2)/0.2/, Y(2)/5.306/ DATA X(3)/0.3/, Y(3)/5.569/, X(4)/0.4/, Y(4)/5.938/ DATA X(5)/0.5/, Y(5)/6.437/, X(6)/0.6/, Y(6)/7.0988/ DATA X(7)/0.7/, Y(7)/7.949/, X(8)/0.8/, Y(8)/9.025/ DATA X(9)/0.9/, Y(9)/10.363/
	DATA MS/1/, MF/4/
100 * *	<pre>WRITE(1,100) FORMAT(//,1H ,20X,'POLYNOMIAL REGRESSION ANALYSIS',/1H ,20X,</pre>
	IF (MF .LE. (N-1)) THEN GO TO 5 ELSE MF = N-1 WRITE(*,*)
	WRITE(*,*) 'DEGREE OF POLYNOMIAL CANNOT EXCEED N-1.' WRITE(*,*) 'REQUESTED MAXIMUM DEGREE TOO LARGE- REDUCED TO',MF ENDIF
5	MFP1 = MF+1 MFP2 = MF+2
	DO 10 I =1,N XN(I) =1.

10 CONTINUE YBAR = 0.0 DO 11 J = 1.N YBAR = YBAR + Y(J)11 CONTINUE YBAR = YBAR/NDO 30 I = 1, MFP1 A(I,1) = 0.0A(I, MFP2) = 0.DO 20 J = 1,N A(I,1) = A(I,1)+XN(J)A(I,MFP2) = A(I,MFP2)+Y(J)*XN(J)XN(J)=XN(J)*X(J) CONTINUE 20 30 CONTINUE DO 50 I = 2,MFP1 A(MFP1,I) = 0.DO 40 J = 1, N $\begin{array}{l} \textbf{A}(\textbf{MFP1},\textbf{I}) = \textbf{A}(\textbf{MFP1},\textbf{I}) + \textbf{XN}(\textbf{J}) \\ \textbf{XN}(\textbf{J}) = \textbf{XN}(\textbf{J}) * \textbf{X}(\textbf{J}) \end{array}$ CONTINUE 40 50 CONTINUE DO 70 J = 2, MFP1 DO 60 I = 1, MF A(I,J) = A(I+1,J-1)60 CONTINUE 70 CONTINUE CALL LUDCMQ (A, MFP1, 10) MSP1 = MS +1 DO 95 I = MSP1,MFP1 DO 90 J = 1,I C(J) = A(J, MFP2)90 CONTINUE CALL SOLNQ (A, C, I, 10) IM1 = I - 1 WRITE (1,110)IM1,(C(J), J=1,I) 110 SSE = 0.0SST = 0.0

```
DO 94 IPT = 1, N
       SUM = 0.0
           DO 93 ICOEF = 2, I
               JCOEF = I - ICOEF + 2
               SUM = (SUM + C(JCOEF)) * X(IPT)
93
            CONTINUE
с
       CALCULATE THE ERROR SUM OF SQUARES AND TOTAL SUM OF SQUARES
           SUM = SUM + C(1)
SSE = SSE + (Y(IPT) - SUM)**2
SST = SST + (Y(IPT) -YBAR)**2
 94
       CONTINUE
с
       CALCULATE THE VARIANCE
       VAR = SSE/(N-I)
       WRITE(1,120)VAR,SSE,SST
       120
с
       CALCULATE CORRELATION OF DETERMINATION AND CORRELATION
ċ
       COEFFICIENT.
       CORD = (1.0 - SSE/SST)
       COR = CORD * *0.5
       125
     *
95
       CONTINUE
       CLOSE (UNIT=1)
       STOP
       ÉND
C
C
C
C
C
        THIS PROGRAM FORMS THE LU EQUIVALENT OF THE SQUARE COEFFICIENT MATRIX A. THE LU, IS THEN RETURNED IN THE
       A MATRIX SPACE. THE UPPER TRIANGULAR MATRIX U HAS ONES
ON ITS DIAGONAL. THESE VALUES ARE NOT INCLUDED IN THE
                         THESE VALUES ARE NOT INCLUDED IN THE RESULT.
С
с
                                                                 ******
       SUBROUTINE LUDCMQ (A, N, NDIM)
```

```
DIMENSION A(1:NDIM, 1:NDIM)
         DO 30 I = 1,N
DO 30 J = 2,N
SUM = 0.0
IF (J .GT. I) THEN
GO TO 15
          ELSE
          JM1 = J-1
          ENDIF
          DO 10 K = 1,JM1
          SUM = SUM + A(I,K) * A(K,J)
          CONTINUE
 10
          A(I,J) = A(I,J) - SUM
          GO TO 30
 15
          IM1 \approx I - 1
          IF (IM1 .EQ. 0.) THEN
          GO TO 25
          ELSE
          DO 20 K = 1, IM1
          SUM = SUM + A(I,K) * A(K,J)
 20
          CONTINUE
          ENDIF
 25
          IF (ABS(A(I,I)) .LT. 1.E-10) THEN GO TO 99
          ELSE
          A(I,J) = (A(I,J) - SUM)/A(I,I)
          ENDIF
 30
          CONTINUE
          RETURN
 99
          WRITE(*,*)'REDUCTION NOT COMPLETED AS SMALL VALUE FOUND '
          WRITE(*,*)'FOR DIVISOR IN ROW',I
          RETURN
          END
С
          THIS PROGRAM FINDS THE SOLUTION TO A SET OF N LINEAR EQUATIONS
THAT CORRESPONDS TO THE RIGHT HAND SIDE VECTOR B.
THE A MATRIX IS THE LU DECOMPOSITION EQUIVALENT TO THE
COEFFICIENT MATRIX OF THE ORIGINAL EQUATIONS PRODUCED BY
LUDMQ. THE SOLUTION VECTOR IS RETURNED IN THE B VECTOR.
00000000
          DO THE REDUCTION STEP.
          *******
         SUBROUTINE SOLNQ ( A, B, N, NDIM)
DIMENSION A(1:NDIM, 1:NDIM), B(1:NDIM)
          B(1) = B(1)/A(1,1)
          DO_{20} I = 2, N
          IM1 = I - 1
          SUM = 0.0
          DO 10 K = 1, IM1
```

SUM = SUM + A(I,K) * B(K)

10	CONTINUE B(I) = (B(I) - SUM) / A(I,I)
20	CONTINUE
с	BACK SUBSTITUTION
	DO 40 J = 2,N NMJP2 = N - J + 2 NMJP1 = N - J + 1 SUM = 0.0
30	DO 30 K = NMJP2, N SUM = SUM + A(NMJP1,K)*B(K) CONTINUE B(NMJP1) = B(NMJP1) - SUM
40	CONTINUE RETURN END

POLYNOMIAL REGRESSION ANALYSIS FOR AN EQUATION TO AN Nth DEGREE : Y=C0+C1.X+C2.X^2+C3.X^3+....+Cn.X^n *********** POLYNOMIAL OF DEGREE 1 COEFFICIENTS ARE: 3.809 6.340 VARIANCE IS: .2982E+00 ERROR SUM OF SQUARES: .2087E+01 TOTAL SUM OF SQUARES: .2620E+02 COEFFICIENT OF DETERMINATION IS: .9203 CORRELATION COEFFICIENT: .9593 POLYNOMIAL OF DEGREE 2 COEFFICIENTS ARE: 5.305 -1.822 8.162 .5950E-02 VARIANCE IS: ERROR SUM OF SQUARES: .3570E-01 TOTAL SUM OF SQUARES: .2620E+02 CORRELATION COEFFICIENT: .9993 POLYNOMIAL OF DEGREE 3 COEFFICIENTS ARE: 4.975 1.339 .659 5.002 VARIANCE IS: .1281E-04 ERROR SUM OF SQUARES: .6405E-04 TOTAL SUM OF SQUARES: .2620E+02 COEFFICIENT OF DETERMINATION IS: 1.0000 CORRELATION COEFFICIENT: 1.0000 ______ POLYNOMIAL OF DEGREE 4 COEFFICIENTS ARE: 4.999 1.011 1.965 3.047 .977 .5557E-06 VARIANCE IS: ERROR SUM OF SQUARES: .2223E-05 TOTAL SUM OF SQUARES: .2620E+02 COEFFICIENT OF DETERMINATION IS: 1.0000 CORRELATION COEFFICIENT: 1.0000

```
PROGRAM PROG14
```

C C C	**************************************
0 0 0 0 0 0 0 0 0 0 0 0 0	AB = COEFFICIENT MATRIX AUGMENTED WITH R.H.S VECTORS. N = NUMBER OF EQUATIONS. NP = TOTAL NUMBER OF COLUMNS IN THE AUGMENTED MATRIX NDIM = FIRST DIMENSION OF MATRIX AB IN THE CALLING PROGRAM. THE SOLUTION VECTORS ARE RETURNED IN THE AUGMENTATION COLUMNS OF A
	DIMENSION A(1:10, 1:11)
	OPEN (UNIT=1, FILE='PRN')
	DATA N/4/ DATA (A(1,J), J=1,5) /51.30, 43.20, 56.40, 47.40, 48.80/ DATA (A(2,J), J=1,5) /11.30, 11.50, 15.50, 8.50, 11.56/ DATA (A(3,J), J=1,5) /29.40, 31.50, 22.50, 30.40, 29.43/ DATA (A(4,J), J=1,5) /8.00, 10.30, 5.60, 13.70, 10.21/
100	WRITE(1,100) FORMAT(1H ,10X,'GAUSSIAN ELIMINATION METHOD FOR A SET OF', $)$
105	WRITE(1,105) FORMAT(' LINEAR EQUATIONS',/1H ,70(1H*),//)
110	WRITE(1,110) FORMAT(30X,'AUGMENTED MATRIX',//)
120	DO 10 I=1,N WRITE(1,120)(A(I,J),J=1,N+1) FORMAT(1X,1P6E12.4)
10	NP =N +1 CONTINUE
	DET=1.0
c c	CALL THE GAUSSIAN ELIMINATION
	CALL GAUSS (A, N, NP, 10, DET)
125	WRITE(1,125)DET FORMAT(//, 10X,'DETERMINANT=', 1X,E12.4)
130	WRITE(1,130) FORMAT(//,30X,'SOLUTION', /1H ,70(1H-)) WRITE(1,140) FORMAT(//,10X//, L , X(L)/, (1H ,70(1H-))
140	FORMAT(//,10X,' I X(I)', /1H ,70(1H-)) DO 20 I =1, N WRITE(1,150) I, A(I,N+1)

```
150
         FORMAT(10X,15,10X,1PE16.6)
         CONTINUE
  20
        WRITE(1,160)
FORMAT(1H,70(1H-))
CLOSE (UNIT=1)
 160
         STOP
         END
с
         SUBROUTINE GAUSS (AB, N, NP, NDIM, DET)
         DIMENSION AB(1:NDIM, 1:NP)
с
         BEGIN THE REDUCTION
         NM1=N-1
         DO 35 I=1,NM1
c
c
         FIND THE ROW NUMBER OF THE PIVOT ROW AND THEN INTERCHANGE
         ROWS INORDER TO PLACE THE PIVOT ELEMENT ON THE DIAGONAL.
         IPVT = I
         IP1 = I+1
         DO 10 \overline{J} \approx IP1, N
         IF (ABS(AB(IPVT,I)) .LT. ABS(AB(J,I))) THEN
         IPVT =J
         ELSE
         ENDIF
 10
         CONTINUE
c
c
         CHECK THAT THE PIVOT ELEMENT IS NOT TOO SMALL. OTHERWISE
         PRINT A MESSAGE AND RETURN
         IF ((ABS(AB(IPVT, I)) .LT. 1.E-5)) THEN
         DET=0.0
         GO TO 99
         ELSEIF( IPVT .EQ. I) THEN
         GO TO 25
         ENDIF
        DO 20 JCOL = I,NP
SAVE=AB(I,JCOL)
             AB(I, JCOL) = AB(IPVT, JCOL)
             AB(IPVT, JCOL)=SAVE
 20
         CONTINUE
         DET=-DET
        REDUCE ALL ELEMENTS BELOW THE DIAGONAL IN THE I-TH ROW.
CHECK FIRST TO SEE IF A ZERO ALREADY PRESENT. IF SO,
SKIP REDUCTION FOR THAT ROW.
с
С
с
         DO 32 JROW = IP1,N
 25
             IF (AB(JROW, I) .EQ. 0.0) THEN
GO TO 32
             ELSE
             RATIO=AB(JROW,I)/AB(I,I)
```

ENDIF DO 30 KCOL = IP1,NP AB(JROW, KCOL) = AB(JROW,KCOL)-RATIO*AB(I,KCOL) 30 CONTINUE 32 35 CONTINUE CONTINUE WE STILL NEED TO CHECK A(N,N) FOR SIZE. с IF (ABS(AB(N,N)). LT. 1.E-5) THEN GO TO 99 ELSE ENDIF BACK SUBSTITUTION с NP1=N+1 DO 50 KCOL =NP1,NP AB(N,KCOL)=AB(N,KCOL)/AB(N,N) DO 45 J= 2,N NVBL = NP1-J L=NVBL+1 VALUE = AB(NVBL,KCOL) DO 40 K= L,N VALUE=VALUE-AB(NVBL,K)*AB(K,KCOL) CONTINUE 40 AB(NVBL, KCOL)=VALUE/AB(NVBL, NVBL) 45 CONTINUE CONTINUE 50 с CALCULATE THE DETERMINANT OF THE MATRIX DO 60 I=1,N DET=DET*AB(I,I) CONTINUE 60 RETURN С MESSAGE FOR A NEAR SINGULAR MATRIX. WRITE(1,100) FORMAT(//,10X,'SOLUTION NOT FEASIBLE. A NEAR ZERO PIVOT WAS',\) 99 100 WRITE(1,110)
FORMAT(' ENCOUNTERED') 110 RETURN END

GAUSSIAN ELIMINATION METHOD FOR A SET OF LINEAR EQUATIONS

AUGMENTED MATRIX

5.1300E+01	4.3200E+01	5.6400E+01	4.7400E+01	4.8800E+01
1.1300E+01	1.1500E+01	1.5500E+01	8.5000E+00	1.1560E+01
2.9400E+01	3.1500E+01	2.2500E+01	3.0400E+01	2.9430E+01
8.0000E+00	1.0300E+01	5.6000E+00	1.3700E+01	1.0210E+01

DETERMINANT= -.1149E+05

	SOLUTION	
I	X(I)	
1 2 3 4	1.172198E-01 3.789395E-01 2.117474E-01 3.053561E-01	

000000000000000000000000000000000000000	GAUSS-SEIDEL ITERATION FOR N SIMULTANEOUS LINEAR EQUATIONS THE ARRAY A CONTAINS THE N × N+1 AUGMENTED COEFFICIENT MATRIX THE VECTOR X CONTAINS THE LATEST APPROXIMATION TO THE SOLUTION THE COEFFICIENT MATRIX SHOULD BE DIAGONALLY DOMINANT. AN INITIAL APPROXIMATION IS SENT TO THE SUBROUTINE IN THE VECTOR X. THE SOLUTION, AS APPROXIMATED BY THE SUBROUTINE IS RETURNED IN X. THE ITERATIONS ARE CONTINUED UNTIL THE MAXIMUM CHANGE IN ANY X COMPONENT IS LESS THAN THE TOLERANCE, TOL. IF THIS CANNOT BE ACCOMPLISHED IN NITER ITERATIONS, A MESSAGE IS PRIMTED. IN TERMINATING THE ITERATIONS, NO ELEMENT OF X MAY UNDERGO A MAGNITUDE CHANGE GREATER THAN TOLERANCE TOL FROM ONE ITERATION TO THE NEXT.
c	PARAMETERS:
000000	 A - AUGMENTED COEFFICIENT MATRIX X - INITIAL APPROXIMATION TO SOLUTION, ALSO RETURNS RESULTS N - NUMBER OF SIMULTANEOUS LINEAR EQUATIONS NITER - LIMIT TO THE NUMBER OF ITERATIONS TOL - TEST VALUE TO STOP THE ITERATING SOLUTION
c	***********
	DIMENSION A(1:20,1:20), X(20)
	INTEGER N, NITER, I, J
	OPEN (UNIT=1, FILE='PRN')
	DATA N, NITER, TOL/4, 15, 0.0001/
	DATA A(1,1)/1100.0/, A(1,2)/0.0/, A(1,3)/0.0/, A(1,4)/0.0/ DATA A(1,5)/1000.0/, A(2,1)/1000.0/, A(2,2)/-1400.0/ DATA A(2,3)/100.0/, A(2,4)/0.0/, A(2,5)/0.0/, A(3,1)/0.0/ DATA A(3,2)/1100.0/, A(3,3)/-1240.0/, A(3,4)/100.0/ DATA A(3,5)/0.0/, A(4,1)/0.0/, A(4,2)/0.0/, A(4,3)/1100.0/ DATA A(4,4)/-1250.0/, A(4,5)/0.0/
с	INITIAL GUESS VALUES
	DATA X(1), X(2), X(3), X(4)/0.5, 0.5, 0.5, 0.5/
	NP1 = N+1
120 125	WRITE(1, 120) FORMAT(5X, 'SOLUTION OF SIMULTANEOUS LINEAR EQUATIONS BY',\) WRITE(1,125) FORMAT(' GAUSS-SEIDEL METHOD', /5X, 70(1H*))
130	WRITE(1,130) N, NITER, TOL FORMAT(//, 10X, 'NUMBER OF LINEAR EQUATIONS:',I4,/,

	+ 10X, 'MAXIMUM ITERATIONS', 14, /, 10X, 'CONVERGING TOLERANCE:', E14.6)	
140	<pre>WRITE(1,140) FORMAT(//,10X,'THE COEFFICIENT MATRIX A(1,1)A(N+1,N+1) IS:',, + 5X, 70(1H-))</pre>	/
	DO 10 I = 1,N WRITE(1,150)(A(I,J),J=1,NP1)	
150	FORMAT(/,5(3X,F12.4),5X)	
10 155	CONTINUE WRITE(1,155) FORMAT(/,5X,70(1H-))	
160	<pre>WRITE(1,160) FORMAT(//,10X,'THE STARTING VECTOR X(1)X(N) IS:')</pre>	
170	WRITE(1,170)(X(I),I=1,N) FORMAT(/,15X,1PE14.6)	
	CALL GSITER (A, X, N, NITER, NP1, TOL)	
	CLOSE (UNIT=1) STOP END	
	SUBROUTINE GSITER (A, X, N, NITER, NP1, TOL) DIMENSION A(1:20, 1:20), X(20) INTEGER N, NITER, FLAG	
с	NORMALIZE DIAGONAL ELEMENTS IN EACH ROW	
	DO 10 I =1, N ASTAR = A(I,I) DO 10 J = 1,NP1 A(I,J) = A(I,J)/ASTAR	
10	CONTINUE	
с	NOW WE PERFORM THE ITERATIONS	
	DO 30 ITER =1, NITER FLAG = 1 DO 40 I = 1,N XSTAR = X(I) X(I) = A(I,NP1)	
с	CALCULATE NEW SOLUTION VALUE, X(I)	
	DO 50 J = 1,N IF (I .EQ. J) THEN GO TO 50 ELSE X(I) = X(I)-A(I,J)*X(J) ENDIF	

50 CONTINUE

```
C TEST X(I) FOR CONVERGENCE

IF (ABS(XSTAR - X(I)) .LE. TOL) THEN

GO TO 40

else

FLAG = 0

ENDIF
```

40 CONTINUE

IF (FLAG .NE. 1) THEN GO TO 30 ELSE

- WRITE(1,100) ITER 100 FORMAT(/,10X,'PROCEDURE CONVERGED AFTER',2X,I2,2X,'ITERATIONS',/)
- WRITE(1,110)(X(I),I =1,N)
 110 FORMAT(/10X,'SOLUTION VECTOR X(1)....X(N) IS:',//,(15X,1PE14.6))
- WRITE (1, 120) 120 FORMAT (5X, 70(1H-))

GO TO 60 ENDIF

- 30 CONTINUE
- 60 RETURN

- WRITE(1,130)TOL,NITER
 130 FORMAT(//,10X,'DID NOT MEET TOLERANCE OF',E14.6,'IN',I4,
 * 'ITERATIONS')
- WRITE(1,140)(X(I),I=1,N)
 140 FORMAT(/,10X,'CURRENT VECTOR X(1).....X(N) IS :',/(15X,1PE14.6))
 RETURN
 END

SOLUTION OF SIMULTANEOUS LINEAR EQUATIONS BY GAUSS-SEIDEL METHOD

NUMBER OF LINEAR EQUATIONS: 4 MAXIMUM ITERATIONS 15 CONVERGING TOLERANCE: .100000E-03

THE COE	FFICIENT MATRI	X A(1,1)A(N+	1,N+1) IS:	
1100.0000	.0000	.0000	.0000	1000.0000
1000.0000	-1400.0000	100.0000	.0000	.0000
.0000	1100.0000	-1240.0000	100.0000	.0000
.0000	.0000	1100.0000	-1250.0000	.0000

THE STARTING VECTOR X(1)....X(N) IS:

```
5.000000E-01
```

5.000000E-01

5.000000E-01

5.000000E-01

PROCEDURE CONVERGED AFTER 5 ITERATIONS

SOLUTION VECTOR X(1)....X(N) IS:

9.090909E-01 6.968780E-01 6.654189E-01

5.855687E-01

PROGRAM PROG16 C C C ***** THIS PROGRAM EMPLOYS THE NEWTON-RAPHSON'S METHOD TO DETERMINE С THE ROOTS OF NONLINEAR EQUATIONS С EXTERNAL FUN.FDERV OPEN (UNIT=1, FILE='PRN') С INPUT DATA FOR INITIAL GUESS OF THE ROOT DATA X/0.015/ WRITE(1,100) 100 FORMAT(15X, 'NEWTON-RAPHSON METHOD FOR A NON-LINEAR EQUATION', /1H ,76(1H*)) WRITE(1,110)X 110 FORMAT(//,6X,'INITIAL GUESS OF THE ROOT:',F12.4) XTOL=1.E-4 FTOL=1.E-5 NLIM=50 I=0 CALL NEWT(FUN, FDERV, X, XTOL, FTOL, NLIM, I) CLOSE (UNIT=1) STOP END с ****************************** c c THIS PROGRAM USES THE NEWTON-RAPHSON'S METHOD TO CALCULATE THE ROOT OF NONLINEAR FUNCTIONS С SUBROUTINE NEWT(FUN, FDERV, X, XTOL, FTOL, NLIM, I) 000000000 FUN = FUNCTION THAT CALCULATES F(X) FOR - FUNCTION THAT CALCULATES T(X)FDERV = FUNCTION THAT CALCULATES THE DERIVATIVE OF FUN X = VALUE OF APPROXIMATE ROOT XTOL,FTOL = TOLERANCE VALUES FOR X, F(X) NLIM = LIMIT TO NUMBER OF ITERATIONS I =0 SHOWS HOW ROUTINE IS TERMINATED =1 MEETS TOLERANCE FOR X VALUES Ι С Ι =2 F(X) 11 11 С Ι с ****** *************************************

LOGICAL PRINT PRINT=.TRUE. IF (I .NE. 0) THEN PRINT=.FALSE.

```
ELSE
     FX=FUN(X)
     ENDIF
     DO 10 J=1,NLIM
     DELTAX=FX/FDERV(X)
     X=X-DELTAX
     FX=FUN(X)
     FDX=FDERV(X)
     IF (.NOT. PRINT) THEN
GO TO 20
     ELSE
ENDIF
   20 IF (ABS(DELTAX) .LE. XTOL) THEN
     GO TO 30
     ELSEIF (ABS(FX) .LE. FTOL) THEN
     GO TO 40
     ELSE
     ENDIF
     CONTINUE
10
с
     WHEN ITERATION IS COMPLETED, NLIM IS EXCEEDED
     T = -1
     WRITE(1,150)NLIM,X,FX
 150 FORMAT(//,1H0,3X,'TOLERANCE NOT MET AFTER',1X,12,1X,
*'ITERATIONS',1X,' THE FINAL ROOT: X=',E10.5,2X,'F(X)=',E10.5)
     RETURN
С
     THIS SECTION RETURNS AFTER MEETING XTOL
  30 I=1
     WRITE(1,160)J,X,FX
 160 FORMAT(//,1H0,3X,'TOLERANCE MET IN',1X,12,1X,
*'ITERATIONS',1X,'THE FINAL ROOT: X=',E10.5,2X,'F(X)=',E10.5)
WRITE (1, 165)
165 FORMAT (1H, 76(1H-))
     RETURN
     THIS SECTION RETURNS AFTER MEETING F(X) TOLERANCE
С
   40 I=2
     WRITE(1,170)J,X,FX
     FORMAT(//,1H0,3X,'TOLERANCE MET IN',1X,12,1X,
*'ITERATIONS',1X,'THE FINAL ROOT: X=',E10.5,2X,'F(X)=',E10.5)
 170
     WRITE (1, 175)
175 FORMAT (1H ,76(1H-))
     RETURN
     END
С
С
      С
     THIS PROGRAM CALCULATES THE VALUE OF THE FUNCTION, FUN
      ċ
```

RETURN END

```
NEWTON-RAPHSON METHOD FOR A NON-LINEAR EQUATION
```

INITIAL GUE:	SS OF	THE ROOT: .0	150	
AT ITERATION	1	X=.19423E-01	F(X)=.22078E+00	
AT ITERATION	2	X≕.20577E-01	F(X)=.93464E-02	
AT ITERATION	3	X=.20630E-01	F(X)=.18002E-04	

TOLERANCE MET IN 3 ITERATIONS THE FINAL ROOT: X=.20630E-01 F(X)=.18002E-04

0000000000000	THIS PROGRAM PERFORMS INTERPOLATION WITHIN A SET OF (X, Y) PAIRS TO GIVE THE Y VALUE CORRESPONDING TO A GIVEN X VALUE. THE PROGRAM USES THE NEWTON-GREGORY FORWARD AND BACKWARD INTER- POLATIONS AND STIRLING'S CENTRAL DIFFERENCE METHOD. THE NUMBER OF DATA AND (X, Y) PAIRS ARE IN A DATA FILE:DATA17.DAT THE RESULTS ARE PRINTED IN A UNIT NUMBER (UNIT=1) AND FILE NAME IS ASSIGNED TO PRINT IN 'PRN'. X = ARRAY OF VALUES OF THE INDEPENDENT VARIABLEY = ARRAY OF VALUES OF THE DEPENDENT VARIABLEXR = THE X - VALUE FOR WHICH THE ESTIMATE OF Y IS REQUIRED
C C C	M = THE NUMBER OF INDEPENDENT VARIABLE TO BE INTERPOLATED N = THE NUMBER OF POINTS **********************************
L.	
	DIMENSION X(1:90),Y(1:90,1:10),XR(1:40) OPEN (UNIT=3,FILE='DATA17.DAT',STATUS='OLD',ERR=18) OPEN (UNIT=1,FILE='PRN') E=0.000005
	READ(3,*,ERR=19)N K=(2*N)-1
18	READ(3,*,ERR=19)(X(I),Y(I,1),I=1,K,2) GO TO 10 WRITE(*,11)
11	FORMAT(3X,'FILE DOES NOT EXIST') GO TO 30
19 22	WRITE(*,22) FORMAT(3X,'ERROR MESSAGE IN THE DATA VALUE') GO TO 30
10	DO J=2,6 N1=K-J
	DO I=J,K,2 Y(I,J)=Y(I+1,J-1)-Y(I-1,J-1) IF (Y(I,J) .GT. E*2**(J-1)) THEN
	K1=J ELSE ENDIF
	END DO END DO
99	WRITE(1,99) FORMAT(30X,'THE DIFFERENCE TABLE',/1H ,70(1H*))
100 110	DO I1=1,3,2 WRITE(1,100)X(I1),(Y(I1,J1),J1=1,I1,2) WRITE(1,110)(Y(I1+1,J1),J1=2,I1+1,2) FORMAT(1H,8X,F8.1,2(8X,F8.4)) FORMAT(1H,32X,2(F8.4,8X)) END DO
	DO I2=5,K-5,2 WRITE(1,200)X(I2),(Y(I2,J2),J2=1,6,2)

```
WRITE(1,250)(Y(I2+1,J2),J2=2,6,2)
 200
         FORMAT(1H ,8X,F8.1,3(8X,F8.4))
FORMAT(1H ,32X,3(F8.4,8X))
 250
         END DO
         DO I3=K-4,K-1,2
         WRITE(1,300) X(I3),(Y(I3,J3),J3=1,K-I3+1,2)
         WRITE(1,350)(Y(13+1,31),J3=2,K-I3,2)
FORMAT(1H, 8X,F8.1,3(8X,F8.4))
FORMAT(1H, 32X,2(F8.4,8X))
 300
 350
         END DO
         WRITE(1,400) X(K),Y(K,1)
 400
         FORMAT(1H ,8X,F8.1,8X,F8.4)
         WRITE(1,450) E
         FORMAT(//,10X,1H ,'ROUND-OFF ERROR:',E12.5,//)
 450
         READ THE NUMBER OF INDEPENDENT VARIABLES TO BE INTERPOLATED
С
         READ(3,*,ERR=19)M
         DO L=1,M
С
         READ THE VALUE OF THE INDEPENDENT VARIABLE
         READ(3,*,ERR=19) XR(L)
         DO I=1,K-1,2
         IF (XR(L) .GT. X(I) .AND. XR(L) .LT. X(I+1)) THEN
         J≃I
         ELSE
         END IF
         END DO
         RC = (XR(L) - X(J)) / ABS(X(3) - X(1))
         R = (XR(L) - X(1)) / ABS(X(3) - X(1))
         T = (XR(L) - X(K)) / ABS(X(3) - X(1))
         CALL NFD(YF,Y,R)
         CALL NBD(YB,Y,T,K)
         CALL SCD(YC,Y,RC,J)
         WRITE(1,540)
        WRITE(1,540)
WRITE(1,550) XR(L),YF
WRITE(1,600) XR(L),YB
WRITE(1,650) XR(L),YC
       FORMAT(20X,1H ,'INTERPOLATED VALUE OF THE INDEPENDENT VARIABLE',
 540
                 /1H ,70(1H*))
 550 FORMAT(/,10X,'NEWTON FORWARD DIFF.',10X,'XR=',F8.2,10X,
*'YF=',F8.4)
600 FORMAT(/,10X,'NEWTON BACKWAD DIFF.',10X,'XR=',F8.2,10X,
     *'YB=',F8.4)
 650 FORMAT(/,10X,'STIRLING CENTRAL DIFF.',8X,'XR=',F8.2,10X,'YC=',
               F8.4)
         END DO
```

700		WRITE (1, 700) FORMAT (1H ,70(1H-))
		CLOSE (UNIT=3,STATUS='KEEP') CLOSE (UNIT=1)
30		STOP END
с с с с с с с с		**************************************
	1	SUBROUTINE NFD(B,C,D) DIMENSION C(1:90,1:10) B=C(1,1)+D*C(2,2)+(D*(D-1.)*C(3,3))/2.0 +(D*(D-1.)*(D-2.)*C(4,4))/6. RETURN END
с с с с		**************************************
	1	SUBROUTINE NED(F,G,H,J) DIMENSION G(1:90,1:10) F=G(J,1)+H*G(J-1,2)+(H*(H+1.)*G(J-2,3))/2.0 +(H*(H+1.)*(H+2.)*G(J-3,4))/6.0 RETURN END
с с с с		**************************************
с	1	SUBROUTINE SCD(P,Q,S,J) DIMENSION Q(1:90,1:10) P=Q(J,1)+(S*((Q(J-1,2)+Q(J+1,2))/2.)/2.)+S*S*Q(J,3) +((S-1.)*S*(S-1.)*((Q(J-1,4)+Q(J+1,4))/2.)/6.) RETURN END

DATA17.DAT						
7 200.0 0.5851 300.0 0.6059 400.0 0.6208 600.0 0.6457 800.0 0.6632 1000.0 0.6735 1200.0 0.6849 1 575.0						
			CE TABLE			
**************************************	********** .5851 .6059 .6208 .6457	.0208 .0149 .0249	0059 .0100 0074	.0159 0174	0333 .0176	.0509
800.0 1000.0	.6632 .6735	.0175	0072 .0011	.0002 .0083	.0081	0095
1200.0	.6849	.0114				
ROUND-OFF ERROR:			MUE THO			12
INTERPOLATED VALUE OF THE INDEPENDENT VARIABLE						
NEWTON FORWARD DI	FF.	XR=	= 575.0	0	¥F=	.6805
NEWTON BACKWAD DI	FF.	XR=	= 575.0	0	YB=	.4388

STIRLING CENTRAL DIFF. XR= 575.00 YC= .6431

```
PROGRAM PROG18
000000000
       ******
       THIS PROGRAM CALCULATES THE AREA OF A FUNCTION DEFINED BY
       THE UNIFORM SPACING OF THE INDEPENDENT X VALUES
       FUN
                   ARRAY OF VALUES OF THE FUNCTION
               =
                   NUMBER OF VALUES
               =
       N
                   THE UNIFORM SPACING BETWEEN X VALUES
       н
               =
       AREA
               =
                  THE ESTIMATED INTERVAL OF THE FUNCTION
с
č
       *****
       DIMENSION F(1:100)
       OPEN (UNIT=3, FILE='DATA18.DAT', STATUS='OLD', ERR=18)
OPEN (UNIT=1, FILE='PRN')
C
C
       READ THE NUMBER OF VALUES AND THE UNIFORM SPACING
       BETWEEN THE X- VALUES
       READ(3,*,ERR=19)N,H
       DO 10 I =1, N
       READ (3,*,ERR=19)F(I)
 10
       CONTINUE
       WRITE(1,100)
 100
       FORMAT(1H ,15X, 'SIMPSON RULE TO CALCULATE THE INTEGRATION', \)
       WRITE(1,105)
 105
       FORMAT(1X,'OF A FUNCTION',/5X,70(1H*))
       GO TO 20
 19
       WRITE(*, 110)
       FORMAT(3X, 'FILE DOES NOT EXIST')
 110
       GO TO 999
       WRITE(*,120)
FORMAT(3X,'ERROR MESSAGE IN THE DATA VALUE')
 18
 120
       GO TO 999
 20
       CALL SIMPS (F, N, H, AREA)
       WRITE(1,130)AREA
       FORMAT(//, 20X, 'AREA OF THE FUNCTION:', F10.4)
 130
       WRITE (1, 140)
 140
       FORMAT (5X,70(1H-))
       CLOSE (UNIT=3, STATUS='KEEP')
CLOSE (UNIT=1)
 999
       STOP
       END
```

```
0000
       THIS PROGRAM PERFORMS SIMPSON'S RULE INTEGRATION OF A
       FUNCTION DEFINED BY A TABLE OF EQUISPACED VALUES.
       SUBROUTINE SIMPS(F, N, H, AREA)
       DIMENSION F(1:100)
       INTEGER N, NP, NH, NB, NE
C
C
       DETERMINE WHETHER NUMBER OF PANELS IS EVEN
       NUMBER OF PANELS IS N-1
       NP=N-1
       NH=NP/2
       NB=1
       AREA=1.0
       IF ((NP-2*NH) .NE. 0) THEN
C
C
C
       NUMBER OF PANELS IS ODD. THEREFORE USE SIMPSON'S 3/8 RULE
       ON FIRST THREE AND 1/3 RULE ON THE REMAINING FUNCTION VALUES.
       AREA=3.0*H/8*(F(1)+3.0*F(2)+3.0*F(3)+F(4))
       NB=4
       ELSEIF (N .EQ. 4) THEN
       RETURN
       ENDIF
С
       USE 1/3 RULE ADD TO FIRST, SECOND AND LAST VALUES.
       AREA=AREA+H/3.0*(F(NB)+4.0*F(NB+1)+F(N))
       NB=NB+2
       IF (NB .EQ. N) THEN
       RETURN
       ENDIF
       NE=N-2
       DO 30 I=NB, NE, 2
       AREA=AREA+H/3.0*(2.0*F(I)+4.0*F(I+1))
 30
       CONTINUE
       RETURN
```

END

DATA18.DAT
22 0.113
0,000
0.308
0.995
0.876 0.786
0.720
0.663
0.606
0.545
0.497
0.450
0.403
0.355
0.313 0.275
0.237
0.213
0.171
0.142
0.123
0.109
0.095
SIMPSON RULE TO CALCULATE THE INTEGRATION OF A FUNCTION

AREA OF THE FUNCTION: 1.0037

с	******	****	*****************		
с	THIS PROGRAM SOLVES A SYSTEM OF N FIRST ORDER DIFFENTIAL				
с	EQUATIONS BY THE RUNGE-KUTTA FOURTH ORDER METHOD.				
č			NS ARE OF THE FORM $DX1/DT = F1(X,T)$		
č			,T), $DX3/DT=F3(X,T)$, ETC.		
č	DAC/DI		, i , j , bk3 / bi i 5 (k) i , j bio.		
č	DERIVS	=	A SUBROUTINE THAT CALCULATES VALUES OF THE		
	DERIVS	-			
С			N DERIVATIVES. IT IS INVOKED BY THE STATEMENT		
с			CALL DERIVS (X, T, F, N)		
с	то	=	INITIAL VALUE OF THE INDEPENDENT VARIABLE.		
с	н	=	THE STEP SIZE, THE INCREMENT TO TO.		
с	X0	=	THE ARRAY THAT HOLDS THE INITIAL VALUES OF THE		
с			FUNCTIONS.		
с	XEND	÷	AN ARRAY THAT RETURNS THE FINAL VALUES OF THE		
с			FUNCTIONS.		
c	XWRK	=	AN ARRAY USED TO HOLD INTERMEDIATE VALUES DURING		
č			CALCULATION. IT MUST BE DIMENSIONED OF SIZE		
c			4 X N IN THE MAIN PROGRAM.		
С	N	=	THE NUMBER OF EQUATIONS IN THE PROGRAM.		
с	F	=	AN ARRAY THAT HOLDS VALUES OF THE DERIVATIVES.		
с	TF	=	FINAL VALUE OF THE INDEPENDENT VARIABLE.		
с	******	****	******************		
	EXTERNA	L DE	RIVS		
	DIMENSI	ON X	O(1:4),F(1:4),XEND(1:4),XWRK(1:4,1:4)		
	REAL K1	. K2	, K3, K4, K5		
	COMMON/				
	COMMON/DATA2/K1,K2,K3,K4,K5 COMMON/DATA3/TF				
	COPINION	DAIA	5/1F		
	ር እምል N ጥድ / ለ 10 0 /				
	DATA N,TF/4, 10.0/				
	OPEN (U	NIT=	1, FILE='PRN')		
	WRITE (1, 1	00)		
100	FORMAT	(15X	, 'RUNGE-KUTTA FOURTH ORDER METHOD FOR A SYSTEM OF')		
	WRITE (i. 1	10)		
110			, 'ORDINARY DIFFERENTIAL EQUATIONS',/,5X, 74(1H*),//)		
		(, , , , , , , , · · (, , , , ,)		
	WRITE (1 1	20)		
120			,'TIME',3X,'CONC. CA',3X,'CONC. CB',3X,'CONC. CC',		
*	TORMAT		, 'CONC. CD',/5X, 74(1H-),/)		
<u>^</u>		27	, CONC. CD ,/SX, /4(In-),/)		
	m o o	~			
	TO = 0.	0			
	H = 0.5				
			40) TO, $(XO(I), I = 1, N)$		
140	FORMAT	(14X	, F8.2, 5X, 4(F6.2, 5X))		
5	CALL RK	SYST	(DERIVS, TO, H, XO, XEND, XWRK, F, N)		
	T0 = T0				
	DO I =				
	X0(1) =				
	END DO	ALI			

```
WRITE (1, 150) TO, (XO(I), I = 1, N)
FORMAT (14X, F8.2, 5X, 4(F6.2, 5X))
 150
         IF (TO .LT. TF) THEN
         GO TO 5
         ELSE
         ENDIF
         WRITE (1, 160)
FORMAT (5X, 74(1H-))
 160
         FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
С
         WRITE (1, *) CHAR(12)
         CLOSE (UNIT = 1)
         STOP
         END
С
с
с
с
с
                                                                           *****
         THIS PROGRAM SOLVES A SYSTEM OF N FIRST ORDER DIFFERENTIAL
         EQUATIONS BY THE RUNGE-KUTTA FOURTH ORDER METHOD.
с
         SUBROUTINE RKSYST (DERIVS, TO, H, XO, XEND, XWRK, F, N)
DIMENSION X0(1:N), XEND(1:N), XWRK(1:4,1:N), F(1:N)
         COMMON/DATA3/TF
         CALCULATE THE FIRST ESTIMATE OF THE DATA X'S
С
         CALL DERIVS (XO, TO, F, N)
         DO I = 1, N
XWRK(1,I) = H * F(I)
             XEND(I) = XO(I) + XWRK(1,I)/2.
         END DO
с
с
         CALCULATE THE SECOND ESTIMATE , THE XEND VECTOR
         HOLDS THE X-VALUES
         CALL DERIVS (XEND, T0+H/2.0, F, N)
         CALL DENT:
DO I = 1, N
XWRK(2,I) = H * F(I)
XWR(2,I) = YO(T) + XWR
             XEND(I) = XO(I) + XWRK(2,I)/2.
         END DO
с
         CALCULATE THE THIRD ESTIMATE
         CALL DERIVS (XEND, T0+H/2.0, F, N)
         DO I = 1, N
XWRK(3,I) = H * F(I)
         XEND(I) = XO(I) + XWRK(3,I)
         END DO
С
         CALCULATE THE FOURTH ESTIMATE
```

```
CALL DERIVS (XEND, TO+H, F, N)
       DO I = 1, N
       XWRK(4,I) = H * F(I)
       END DO
       COMPUTE THE X VALUES AT THE END OF THE INTERVAL FROM A WEIGHTED AVERAGE OF THE FOURTH ESTIMATES.
С
č
       DO I=1,N
       XEND(I)=XO(I)+(XWRK(1,I)+2.0*XWRK(2,I)+2.*XWRK(3,I)+
    + XWRK(4,I))/6.
       END DO
       RETURN
       END
       С
c
c
       THIS PROGRAM DEFINES THE FUNCTIONS OF THE ORDINARY DIFFERENTIAL
       EQUATIONS IN TERMS OF THE F'S AND THE X'S.
С
       SUBROUTINE DERIVS (X, T, F, N)
       DIMENSION X(1:N), F(1:N)
       COMMON/DATA2/K1,K2,K3,K4,K5
       REAL K1, K2, K3, K4, K5
       F(1) = K5 * X(2) - K1 * X(1)
       F(2) = K1 * X(1) + K4 * X(4) - (K2 + K3 + K5) * X(2)
       F(3) = K2 * X(2)
       F(4) = K3 \times X(2) - K4 \times X(4)
       RETURN
       END
0
0
0
0
0
       THIS STORES THE DATA FOR THE RATE CONSTANTS AND THE INITIAL
       STARTING VALUES OF THE CONCENTRATIONS.
       *****
      BLOCK DATA
       DIMENSION X0(4)
       COMMON/DATA1/X0
       COMMON/DATA2/K1,K2,K3,K4,K5
       REAL K1,K2,K3,K4,K5
       DATA X0/9.9,0.0,0.0,0.5/
       DATA K1/0.4/,K2/0.16/,K3/0.13/,K4/0.08/,K5/0.10/
       END
```

CHAPTER 2

Physical Property of Liquids and Gases

INTRODUCTION

Physical property data of compounds are frequently required for pilot plant studies or for the design of large process plants. These data are not readily available except in some simulation design packages or in the largest libraries. The data can, of course, be obtained by conducting experiments to measure the properties of individual substances or of mixtures, which may exhibit nonideal behavior. However this is often time-consuming and expensive. Therefore engineers may resort to correlations and interpolations to obtain data of sufficient accuracy for design purposes. Thermodynamic data of this type is required in most calculations such as sizing vessels, process pipe lines, separation of multicomponents, gas absorption and chemical reactor design.

Many predictions and correlations of thermodynamic physical data have been published [1-3]. This chapter reviews physical property data for liquids and gases and presents computer programs to calculate these properties for a range of temperature and correlation constants. Appendix A provides tables of some selected physical property data.

Physical Property Data of Water

The programs described cover thermodynamic data of water over a range of temperature from 0°C to 350°C. These data are for density, viscosity, heat capacity, thermal conductivity, surface tension, latent heat of vaporization, and vapor pressure.

Density of Liquids

Saturated liquid densities at any temperature are based on the correlation [1]:

$$\rho_{\rm L} = AB^{-(1-T_{\rm r})^{2/7}} \tag{2-1}$$

where ρ_L = saturated liquid density, g/cm³.

A and B = correlation constants for a chemical compound.

 T_r = reduced temperature, T/T_c .

Many methods [4,5,6] have been proposed to estimate specific volumes or densities of pure liquids. All are based on the law of corresponding states. They are complex algebraically and therefore require the aid of a computer when many calculations are involved. Appendix A1 gives tabulated values of the correlation constants A and B for some chemical compounds. Computer program PROG21 employs Equation 2-1 for estimating the density of liquids. Typical results for water are illustrated in Table 2-1. Figure 2-1 shows the density of water over the range from 0°C to a critical value of $T_c = 374.2^{\circ}C$.

Viscosity of Liquids

The viscosity of a saturated liquid as a function of temperature can be expressed by,

$$\log \mu_{L} = A + \frac{B}{T} + CT + DT^{2}$$
(2-2)

where μ_L = viscosity of saturated liquid, cP A, B, C, and D = correlation constants T = temperature, K

The viscosities of liquids decrease with temperature. Also, the variation is linear over a wide range of temperatures from the freezing point to the boiling point. This is expressed by the Andrade correlation [7],

$$\ln \mu_{\rm L} = \ln A + \frac{B}{T} \tag{2-3}$$

Equation 2-3 is used to correlate the effect of temperature on liquid viscosity. However, it fails at low temperatures because liquids show a

Table 2-1
Computer Output of Physical Properties of Water

	1	TEMPERATURE oC	
	2	25.00	
	3	50.00	
	4	100.00	
	5	150.00	
	6 7	200.00	
	8	250.00 300.00	
	9	350.00	
	10	374.20	
******		TER: 0oC - 374.2oC	*****
	TEMPERATURE oC	DENSITY g/cm.^3	
1	.0	1.267	
2	25.0	1.235	
ŝ	50.0	1.203	
4	100.0	1.135	
5	150.0	1.062	
6	200.0	.983	
7	250.0	.893	
8 9	300.0	.784	
10	350.0 374.2	.627 .095	
*****		MATER: 00C - 374.20C	*****
	TEMPERATURE oC	VISCOSITY CP	
	.0	1.732	
1 2	.0 25.0	1.732	
		1.732 .911 .561	
2 3 4	25.0	.911	
2 3 4 5	25.0 50.0 100.0 150.0	.911 .561	
2 3 4 5 6	25.0 50.0 100.0 150.0 200.0	.911 .561 .291 .192 .142	
2 3 4 5 6 7	25.0 50.0 100.0 150.0 200.0 250.0	.911 .561 .291 .192 .142 .109	
2 3 5 6 7 8	25.0 50.0 100.0 150.0 200.0 250.0 300.0	.911 .561 .291 .192 .142 .109 .082	
2 3 4 5 6 7	25.0 50.0 100.0 150.0 200.0 250.0	.911 .561 .291 .192 .142 .109	

(continued)

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.

Table 2-1 (continued)

	TEMPERATURE oc	HEAT CAPACITY cal/gm.oC
1	.0	.996
2	25.0	1.000
3	50.0	1.003
4	100.0	1.010
5	150.0	1.022
6	200.0	1.048
7	250.0	1.092
8	300.0 350.0	1.163 1.265
*****	THERMAL CON	DUCTIVITY OF WATER: 0oC - 350oC
	TEMPERATURE OC	THERMAL CONDUCTIVITY micro-cal/s.cm.K
1	.0	1375.736
2	25.0	1472.146
3	50.0	1549.540
4	100.0	1647.284
5	150.0	1668.968
6	200.0	1614.592
7	250.0	1484.156
8	300.0	1277.660
9	350.0	995.104
******		NSION OF WATER: 0oC - 374.2oC
	TEMPERATURE oC	SURFACE TENSION:dyne/cm.
		76.110
1	.0	76.119
2	25.0	71.970
3	50.0	67.765
4	100.0	58.910
5	150.0	46.557
6	200.0	34.664
7	250.0	23.341
8	300.0	12.782
9	350.0	3.450
10	374.2	.000
****		SURE OF WATER: 0oC - 374.2 oC
	TEMPERATURE oC	VAPOR PRESSURE: mm Hg
1	.0	4.593

106

•

Table 2-1 (continued)

25.0 50.0 100.0 250.0 250.0 300.0 350.0 374.2 HEAT-OF-VAPC	23.787 92.665 761.613 3578.623 11679.780 29846.610 64535.430 124430.400 165952.400
100.0 150.0 200.0 250.0 300.0 350.0 374.2	92.665 761.613 3578.623 11679.780 29846.610 64535.430 124430.400 165952.400
150.0 200.0 250.0 300.0 350.0 374.2	3578.623 11679.780 29846.610 64535.430 124430.400 165952.400
200.0 250.0 300.0 350.0 374.2	11679.780 29846.610 64535.430 124430.400 165952.400
250.0 300.0 350.0 374.2	29846.610 64535.430 124430.400 165952.400
300.0 350.0 374.2	64535.430 124430.400 165952.400
350.0 374.2	124430.400 165952.400
374.2	165952.400
HEAT-OF-VAPO	DRIZATION OF WATER:00C-374.2 oC
*************	*********
EMPERATURE oC	HEAT-OF-VAPORIZATION:cal/g
.0	606.263
25.0	590.540
	574.104
100.0	538.700
150.0	499.026
200.0	453.398
200.0 250.0	453.398 398.701
200.0	453.398

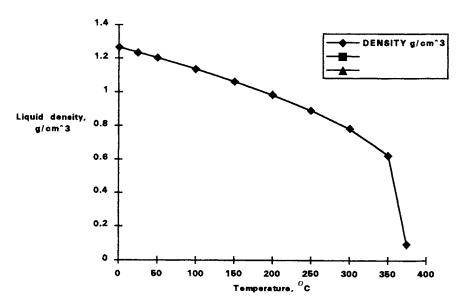


Figure 2-1. Liquid density of water as a function of temperature.

(text continued from page 104)

sharp increase in viscosity as the freezing point is approached. There are other methods in estimating liquid viscosities of compounds [2]. Figure 2-2 gives a plot of liquid viscosity of water from 0°C to its critical value of 374.2°C. Program Prog21 gives a routine for calculating the viscosity of liquids. Table 2-1 gives the viscosity of water as a function of temperature.

Heat Capacity of Liquids

Heat capacity data for liquids are relevant in process design of heat exchanger equipment and chemical reactors. For example, heat capacities are needed in the design of liquid-phase chemical reactors that involve energy input for heating liquid reactants to the reaction temperature. Alternatively, they are essential when sizing reboilers and condensers for distillation plant and when rating of heat exchangers in general.

Correlation constants for liquid heat capacity are expressed in a polynomial of the form:

$$C_{p} = A + BT + CT^{2} + DT^{3}$$
 (2-4)

where C_p = heat capacity of saturated liquid, cal/g°C A, B, C, and D = correlation constants for a chemical compound T = temperature, K

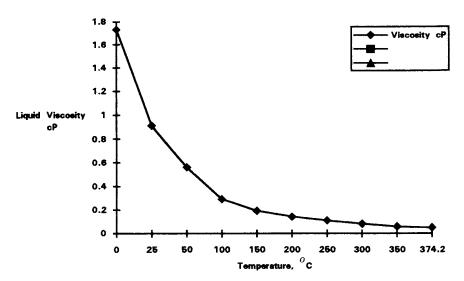


Figure 2-2. Liquid viscosity of water as a function of temperature.

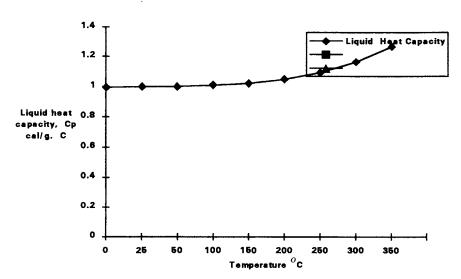
Liquid heat capacities are not strongly dependent on temperatures, except at reduced temperature, $(T_r = T/T_c)$, where $T_r = 0.7$ to 0.8. At high reduced temperatures, liquid heat capacities are large and strongly dependent on temperature. At the boiling point of most organic compounds, heat capacities are between 0.4 and 0.5 cal/g.K. However, in this temperature range, there is no effect of pressure [2]. The computer program PROG21 uses Equations 2-4 for estimating the heat capacity of liquids. Table 2-1 shows the heat capacity of water as a function of temperature. Figure 2-3 shows the heat capacity of water from 0°C to 350°C.

Thermal Conductivity of Liquids

Liquid thermal conductivities, k_L , are required in many chemical and process engineering applications where heat transfer is prevalent. They are required to evaluate the Nusselt number hd/k, the Prandtl number $c\mu/k$, and in correlations to predict the idealized condensing film coefficient based upon laminar liquid flow over a cooled surface. The thermal conductivity of a saturated liquid is:

$$\mathbf{k}_1 = \mathbf{A} + \mathbf{B}\mathbf{T} + \mathbf{C}\mathbf{T}^2 \tag{2-5}$$

where k_L = thermal conductivity of saturated liquid, microcal/s.cm.°C A, B, and C = correlation constants



T = temperature, K

Figure 2-3. Liquid heat capacity of water as a function of temperature.

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Values of k_L for most common organic liquids range between 250 and 400 µcal/cm.s.°C at temperatures below the normal boiling point. Water and other highly polar molecules have values that are two to three times larger. Except for water, aqueous solutions, and multihydroxy molecules, the thermal conductivity of most liquids decreases with temperature. Below or close to the normal boiling point, the decrease is nearly linear. Methods of computing k_L have been reviewed [2]. The computer program PROG21 gives a routine for calculating k_L of liquids, and Table 2-1 shows the results for water. Figure 2-4 is a plot of thermal conductivity of water from 0°C to 350°C.

Surface Tensions of Liquids

Surface tension data of liquids are important in many process design calculations for situations where these is a two-phase interface, e.g., two-phase flow, distillation, absorption and condensation. Surface tension can be expressed as:

$$\sigma = \sigma_1 \left[\frac{T_c - T}{T_c - T_1} \right]^R$$
(2-6)

where σ_1 = surface tension at $T_1^{\circ}C$, dynes/cm T_c = critical temperature, K

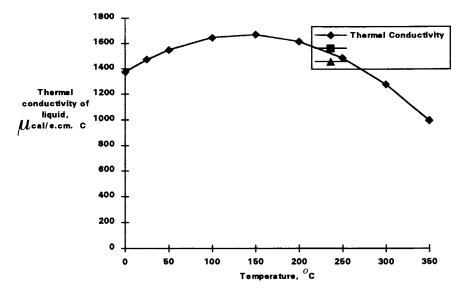


Figure 2-4. Thermal conductivity of water as a function of temperature.

T = temperature, K R = correlation parameter

For water, the temperature range for which surface tension is valid is:

 $\begin{array}{ll} \text{Temperature range 0°C to 100°C} & \sigma_1 = 71.97 \\ \text{Temperature range 100°C to 374.2°C} & \sigma_1 = 58.91 \\ \text{T}_1 = 298.16 \text{ K}, \text{ R} = 0.8105 \text{ for 0°C to 100°C} \\ \text{T}_1 = 373.16 \text{ K}, \text{ R} = 1.1690 \text{ for 100°C to 374.2°C}. \end{array}$

The surface tension has been correlated with other physical parameters such as liquid compressibility, viscosity, molar fractions, and the refractive index. Rao *et al.* [8] developed a linear relationship between the surface tension at normal boiling point (log σ_b) and the reduced boiling point temperature (T_{br}). Hadden [9] presented a nomograph for hydrocarbons that enables rapid calculation of σ . For cryogenic liquids, Sprows and Prausnitz [10] introduced the equation

 $\sigma = \sigma_0 (1 - T_r)^p \tag{2-7}$

where σ = surface tension, dyne/cm

 σ_0 and p were determined by a least-squares analysis of the measured data. The computer program PROG21 gives a routine for estimating the surface tension of liquids, and typical results for water are shown in Table 2-1. Figure 2-5 demonstrates the surface tension of water from 0°C to its critical value of 374.2°C. In general, the surface tension of a liquid in equilibrium with its own vapor decreases with temperature and becomes zero at the critical point. In the reduced temperature range 0.45 to 0.65, σ for most organic liquids range from 20 to 40 dyne/cm. For water, $\sigma = 71.97$ dyne/cm at 25°C.

Vapor Pressure

Vapor pressure data, P_v , of pure components are important in calculations relating to vapor-liquid phase equilibria, e.g., in the simplest case to predict the pressure in a closed vessel containing a specific liquid or mixture of liquids. P_v data are required for bubble point and most dew point computations. These values are used in flash calculations involving mass transfer operations. Clearly the design of pressure requirements for storage equipment requires knowledge of the vapor pressure of the components as does the design of appropriate pressure relief

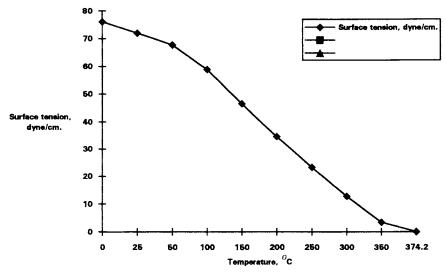


Figure 2-5. Surface tension of water as a function of temperature.

systems. The vapor pressure of the saturated liquid as a function of temperature is

$$\log P_v = A + \frac{B}{T} + C \log T + DT + ET^2$$
 (2-8)

where $P_v =$ vapor pressure of the saturated liquid, mm Hg A, B, C, D, and E = correlation constants for the chemical compound

In the table of vapor pressure, there are values for the acentric factor ω , which is:

$$\omega = -\log P_r - 1.0 \text{ at } T_r = 0.7 \tag{2-9}$$

where reduced pressure $P_r = P/P_c$ reduced temperature $T_r = T/T_c$

The acentric factor is used in thermodynamic correlations involving fugacity, compressibility factor, enthalpy, fugacity, and virial coefficients. The computer program PROG21 provides a routine for estimating the vapor pressure, and Table 2-1 shows P_v of water as a function of temperature. Figure 2-6 shows the vapor pressure of water as a function of temperature to its critical value of 374.2°C. For water, deviations of less

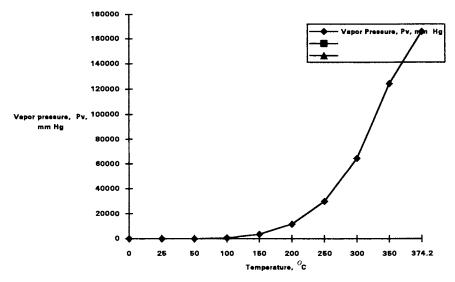


Figure 2-6. Vapor pressure of water as a function of temperature.

than 0.5% are experienced among the other methods. An equation often used to estimate the vapor pressure is the Antoine [11] equation.

$$\ln P_{vp} = A - \frac{B}{T+C}$$
(2-10)

The constants A, B, and C for many compounds are in the data bank of some simulation design packages. Alternatively, these are found in texts such as Reid *et al* [2]. Equation 2-10 is only applicable for pressures with ranges from 10 to 1500 mmHg. Reid *et al* [2]. and Beaton and Hewitt [3] have discussed other methods in estimating vapor pressure of compounds but none appears to offer any specific advantage.

Enthaply of Vaporization

The enthalpy of vaporization, ΔH_v , is also termed the latent heat of vaporization. ΔH_v is the difference between the enthalpy of the saturated vapor and that of the saturated liquid at the same temperature. The enthalpy of vaporization data is used in process calculations such as the design of relief systems involving volatile compounds. In distillation, heat of vaporization values are needed to find the heat loads for the reboiler and condenser. Watson [12] has expressed a widely used correlation in calculating ΔH_v .

$$\Delta H_{v} = \Delta H_{v1} \left[\frac{T_{c} - T}{T_{c} - T_{1}} \right]^{R}$$
(2-11)

where $\Delta H_v =$ heat of vaporization at any temperature cal/g $\Delta H_{vi} =$ heat of vaporization at T₁, cal/g T_c = temperature, K R = characteristic constant for the chemical = 0.38

 ΔH_v decreases with temperature and is zero at the critical point. The computer program PROG21 incorporates a routine for estimating ΔH_v , and Table 2-1 shows the results of water as a function of temperature. Figure 2-7 shows the heat of vaporization of water as a function of temperature. The shape of this plot agrees with most other enthalpy of vaporization data.

Heat Capacities of Gases

The heat capacity of gases is essential for some process engineering design, e.g., for air-cooled heat exchanger and for gas-phase chemical reactions. Here, in the latter case the heat capacities, C_p^0 , for gases are required to determine the heat necessary to bring the chemical compound increase to the reaction temperature. C_p^0 is also used in the rating

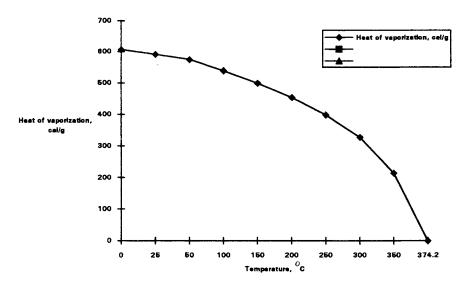


Figure 2-7. Heat of vaporization of water as a function of temperature.

of heat exchangers and energy balance computations. The heat capacity of a mixture of gases may be found from the heat capacities of the individual components contained in the mixtures.

The correlation for C_p^0 of the ideal gas at low pressure is a thirddegree polynomial, which is a function of temperature.

$$C_{n}^{0} = A + BT + CT^{2} + DT^{3}$$
 (2-12)

where C_p^0 = heat capacity of ideal gas at low pressure, cal/gmol.K A, B, C, and D = constants for the chemical compounds T = temperature, K

The correlation coefficients A, B, C, and D can be determined using a least square fit of the experimental data, as discussed in Chapter 1. Huang and Daubert [13] published constants for a fourth-order polynomial for heat capacities of many hydrocarbons. Yuan and Mok [14] used an exponential temperature function to correlate C_p^0 with temperature and provided constants for many hydrocarbons and nonhydrocarbons. Tans [15] produced a nomograph that allows a rapid approximation of C_p^0 for paraffins. A compedium of C_p^0 as a function of temperature for several hundreds of organic compounds is given by Stull *et al.* [16] and in JANAF [17] tables. PROG22 provides a routine for estimating C_p^0 of gases, and Table 2-2 shows results of carbon dioxide. Figure 2-8 is a plot of C_p^0 with temperature for carbon dioxide.

Heats of Formation

Heats of formation, ΔH_r^0 , for individual chemicals involved in chemical reactions are important to find the heat of reaction, ΔH_r^0 , and associated heating and cooling requirements. If $\Delta H_r^0 < 0$, then the chemical reaction is exothermic and will require cooling. If $\Delta H_r^0 > 0$, the reaction is endothermic and heating is therefore required.

The correlation of ΔH_f^0 of the ideal gas at low temperature is based on a series expansion in temperature and is expressed as:

$$\Delta H_f^0 = A + BT + CT^2 \tag{2-13}$$

where ΔH_f^0 = heat of formation of ideal gas at low pressure, kcal/g-mol A, B, and C = corellation constants

T = temperature, K

****		ROPERTY DATA OF GASES
		TEMPERATURE RANGE (300 - 1500K)
	1	300.00
	2	400.00
	3	500.00
	4	600.00
	5	700.00
	6	800.00
	7	900.00
	8	1000.00
	9	1100.00
	10	1200.00
	11	1300.00
	12	1400.00
	13	1500.00
		RMATION OF METHANE

	TEMPERATURE K	HEAT OF FORMATION kcal/mol
1	300.0	-17.962
2	400.0	-18.676
3	500.0	-19.320
4	600.0	-19.894
5	700.0	-20.398
6	800.0	-20.832
7	900.0	-21.196
8	1000.0	-21.490
9	1100.0	-21.714
10	1200.0	-21.868
11	1300.0	-21.952
12 13	1400.0 1500.0	-21.966 -21.910
*****		NERGY OF FORMATION OF ETHYLENE kcal/mol.
	TEMPERATURE K	FREE ENERGY OF FORMATION:kcal/mol
1	300.0	15.830
2	400.0	17.640
3	500.0	19.450
	600.0	21.260
4		
4 5	700.0	23.070

Table 2-2Computer Output of Physical Properties of Gases

Table 2-2 (continued)

8	1000.0	28.500	
9	1100.0	30.310	
10	1200.0	32.120	
11	1300.0	33.930	
12	1400.0	35.740	
13	1500.0	37.550	
******		TTY OF CARBON DIOXIDE	* * * * * * * * * * * * * * * * *
	TEMPERATURE K	HEAT CAPACITY cal/gm-mol.	к
1	300.0	9.201	
2	400.0	10.134	
3	500.0	10.927	
4	600.0	11.594	
5	700.0	12.149	
6 7	800.0 900.0	12.607 12.983	
8	1000.0	13.290	
9	1100.0	13.544	
10	1200.0	13.758	
11	1300.0	13.948	
12	1400.0	14.128	
13	1500.0	14.312	
*****		DUCTIVITY OF PROPANE:26.80C	*******
*****	***************	*********************	************
***************************************	TEMPERATURE oC	THERMAL CONDUCTIVITY mic	************
2	26.8 126.8	42.432 70.267	************
2 3	TEMPERATURE oC	THERMAL CONDUCTIVITY mic	************
2	26.8 226.8 226.8	42.432 70.267 103.678	*******
2 3 4 5 6	26.8 126.8 226.8 326.8 326.8 426.8 526.8	42.432 70.267 103.678 141.461 182.412 225.324	************
2 3 4 5 6 7	26.8 126.8 226.8 326.8 326.8 326.8 426.8 526.8 526.8 626.8	42.432 70.267 103.678 141.461 182.412 225.324 268.995	************
2 3 4 5 6 7 8	26.8 126.8 226.8 326.8 326.8 426.8 526.8 626.8 626.8 726.8	42.432 70.267 103.678 141.461 182.412 225.324 268.995 312.218	************
2 3 4 5 6 7 8 9	26.8 126.8 226.8 226.8 326.8 426.8 526.8 526.8 626.8 726.8 826.8	42.432 70.267 103.678 141.461 182.412 225.324 268.995 312.218 353.789	*******
2 3 4 5 6 7 8	26.8 126.8 226.8 326.8 326.8 426.8 526.8 626.8 626.8 726.8	42.432 70.267 103.678 141.461 182.412 225.324 268.995 312.218	*******
2 3 4 5 6 7 8 9 10	26.8 126.8 226.8 326.8 426.8 526.8 626.8 626.8 726.8 826.8 826.8 926.8	42.432 70.267 103.678 141.461 182.412 225.324 268.995 312.218 353.789 392.504	*******
2 3 4 5 6 7 8 9 10 11	TEMPERATURE OC 26.8 126.8 226.8 326.8 426.8 526.8 626.8 726.8 826.8 926.8 1026.8 1026.8 1026.8 1026.8	42.432 70.267 103.678 141.461 182.412 225.324 268.995 312.218 353.789 392.504 427.156	**************************************
2 3 4 5 6 7 8 9 10 11	TEMPERATURE OC 26.8 126.8 226.8 326.8 426.8 526.8 626.8 726.8 826.8 926.8 1026.8 1026.8 1026.8 1026.8	42.432 70.267 103.678 141.461 182.412 225.324 268.995 312.218 353.789 392.504 427.156	**************************************
2 3 4 5 6 7 8 9 10 11	TEMPERATURE OC 26.8 126.8 226.8 326.8 426.8 526.8 626.8 726.8 8226.8 926.8 1026.8 THE VISCOS TEMPERATURE OC	42.432 70.267 103.678 141.461 182.412 225.324 268.995 312.218 353.789 392.504 427.156	**************************************
2 3 4 5 6 7 8 9 10 11	TEMPERATURE oC 26.8 126.8 226.8 326.8 426.8 526.8 626.8 726.8 826.8 926.8 1026.8 THE VISCOS TEMPERATURE oC 26.8	42.432 70.267 103.678 141.461 182.412 225.324 268.995 312.218 353.789 392.504 427.156	**************************************
2 3 4 5 6 7 8 9 10 11 11	**************************************	<pre>************************************</pre>	**************************************
2 3 4 5 6 7 8 9 10 11 11	TEMPERATURE OC 26.8 126.8 226.8 326.8 426.8 526.8 626.8 726.8 826.8 926.8 1026.8 THE VISCOS THE VISCOS TEMPERATURE OC 26.8 126.8 226.8	42.432 70.267 103.678 141.461 182.412 225.324 268.995 312.218 353.789 392.504 427.156	**************************************
2 3 4 5 6 7 8 9 10 11 	**************************************	<pre>************************************</pre>	**************************************
2 3 4 5 6 7 8 9 10 11 	**************************************	<pre>************************************</pre>	**************************************
2 3 4 5 6 7 8 9 10 11 	TEMPERATURE oC 26.8 126.8 226.8 326.8 426.8 526.8 626.8 726.8 826.8 926.8 1026.8 1026.8 TEMPERATURE oC 26.8 126.8 126.8 226.8 126.8 126.8 226.8 126.8	42.432 70.267 103.678 141.461 182.412 225.324 268.995 312.218 353.789 392.504 427.156 ITY OF ETHANE: 26.80C-1026. VISCOSITY CP. .009 .012 .015 .017 .019 .022 .024	**************************************
2 3 4 5 6 7 8 9 10 11 11 	TEMPERATURE oC 26.8 126.8 226.8 326.8 426.8 526.8 626.8 726.8 826.8 926.8 1026.8 THE VISCOS TEMPERATURE oC 26.8 1	42.432 70.267 103.678 141.461 182.412 225.324 268.995 312.218 353.789 392.504 427.156 .009 .012 .015 .019 .022 .024 .026	**************************************
2 3 4 5 6 7 8 9 10 11 11 	TEMPERATURE OC 26.8 126.8 226.8 326.8 426.8 526.8 626.8 926.8 1026.8 THE VISCOS THE VISCOS TEMPERATURE OC 26.8 126.8 226.8 326.8 126.8 226.8 126	THERMAL CONDUCTIVITY mic 42.432 70.267 103.678 141.461 182.412 225.324 268.995 312.218 353.789 392.504 427.156 ITY OF ETHANE: 26.80C-1026. VISCOSITY CP. .009 .015 .017 .019 .022 .024 .028	**************************************
2 3 4 5 6 7 8 9 10 11 11 	TEMPERATURE oC 26.8 126.8 226.8 326.8 426.8 526.8 626.8 726.8 826.8 926.8 1026.8 THE VISCOS TEMPERATURE oC 26.8 1	42.432 70.267 103.678 141.461 182.412 225.324 268.995 312.218 353.789 392.504 427.156 .009 .012 .015 .019 .022 .024 .026	**************************************

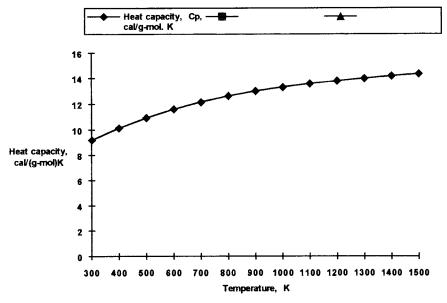


Figure 2-8. Heat capacity of carbon dioxide as a function of temperature.

(text continued from page 115)

Heats of formation are also found from measured heats of combustion, ΔH_c^0 , or from an experimental enthalpy of reaction, ΔH_r^0 . Domalski [18] reviewed values of $\Delta H_{f,298K}^0$ and $\Delta H_{c,298K}^0$ for several hundreds of organic compounds. The computer program PROG22 employs Equation 2-13 in a routine for estimating ΔH_f^0 as a function of temperature. Typical results for methane is presented in Table 2-2. Figure 2-9 shows a profile of ΔH_f^0 with temperature for methane between 298K and 1500K.

Free Energies of Formation

Few methods have been used to estimate the Gibbs free energy ΔG_f^0 directly. Often, ΔH_f^0 and the entropy of formation, ΔS_f^0 , are determined separately. Equation 2-14 can be used to determine ΔG_f^0 .

$$\Delta G_{f}^{0} = \Delta H_{f}^{0} - T\Delta S_{f}^{0}$$
(2-14)

 ΔG_{f}^{0} is often important in chemical reaction engineering. Values of ΔG_{f}^{0} for individual chemicals are required to estimate the free energy of reaction, ΔG_{r}^{0} , and the corresponding chemical thermodynamic equilibrium for the reaction. Also, the following criteria are generally true for the feasibility of chemical reactions. A reaction is considered as promising, when

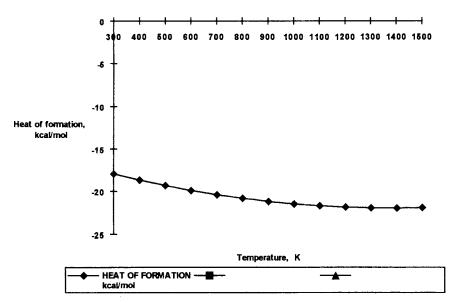


Figure 2-9. Heat of formation of methane as a function of temperature.

 $\Delta G_r^0 < 0$ kcal/gmol

It may possibly be promising, if

 $0 < \Delta G_r^0 < 10$ kcal/gmol

and not promising, if

 $\Delta G_{f}^{0} > 10$ kcal/gmol

Correlation constants for Gibbs free energy of formation of the ideal gas at low pressure are based on a linear relationship in temperature as:

$$\Delta G_f^0 = A + BT \tag{2-15}$$

where ΔG_f^0 = free energy of formation of ideal gas at low pressure, kcal/gmol A and B = correlation constants

T = temperature, K

A routine for calculating ΔG_f^0 is in PROG22, and typical results for ethylene are in Table 2-2. Figure 2-10 shows a plot of free energy for ethylene with temperature.

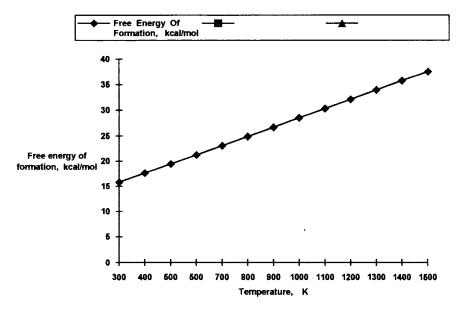


Figure 2-10. Free energy of formation of ethylene as a function of temperature.

Thermal Conductivity of Gases

The thermal conductivity, k_{c} of low pressure gases increases with temperature. In small temperature ranges, k_G, represents some form of linear relation. However, over wide temperature ranges, k_G increases significantly more rapidly with temperature than implied by a linear function. Gas thermal conductivity data are used, e.g., in the design of process equipment and unit operations involving heat transfer, such as the rating of heat exchangers and fluidized bed operations handling gases. Thermal conductivity of a gas can be correlated as a function of temperature as:

$$k_{G} + A + BT + CT^{2} + DT^{3}$$
 (2-16)

where

 k_{G} = thermal conductivity of the gas at low pressure µcal/cm.s.K A, B, C, and D = correlation constantsT = temperature, K

Figure 2-11 shows k_G of propane as a function of temperature. Reid et al. [2] have reviewed other methods for estimating k_{g} .

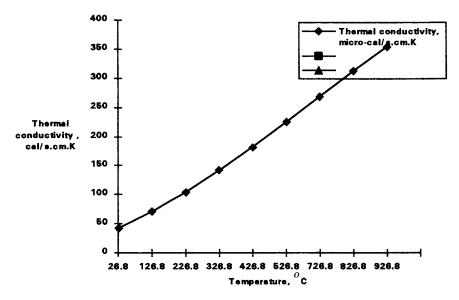


Figure 2-11. Thermal conductivity of propane as a function of temperature.

Viscosity of Gases

Gas phase viscosity data, μ_G , are used in the design of compressible fluid flow and unit operations. For example, the viscosity of a gas is required to determine the maximum permissible flow through a given process pipe size. Alternatively, the pressure loss of a given flowrate can be calculated. Viscosity data are needed for the design of process equipment involving heat, momentum, and mass transfer operations. The gas viscosity of mixtures is obtained from data for the individual components in the mixture.

The correlation for the viscosity of the gas at low pressure can be expressed as:

$$\mu_{\rm c} = (A + BT + CT^2) \times (1 \times 10^{-4})$$
(2-17)

where μ_G = viscosity of the gas at low pressure, cP A, B, and C = correlation constants T = temperature, K

The Herning-Zipperer [19] correlation for the calculation of gas mixture viscosity at low pressure is 122 Fortran Programs for Chemical Process Design

$$\mu_{m} = \frac{\sum MF_{i} \mu_{i} (MW_{i})^{0.5}}{\sum MF_{i} (MW_{i})^{0.5}}$$
(2-18)

where μ_m = gas mixture viscosity, cP MF_i = mole fraction of component i μ_i = gas viscosity of component i MW_i = molecular weight of component i

Reid *et al.* [2] have reviewed other correlation methods for determining μ_G at low pressure. The computer program PROG22 gives a subroutine for estimating gas viscosity and Figure 2-12 gives the viscosity of ethane as a function of temperature. Table 2-2 gives results of viscosity of ethane with temperature.

DIFFUSION COEFFICIENTS (DIFFUSIVITIES)

Diffusion coefficients are important in the design of mass transfer operations equipment, such as gas absorption, distillation and liquidliquid extraction. Experimental data for the common systems can be found in the literature, but for most design work, the values will have to be estimated.

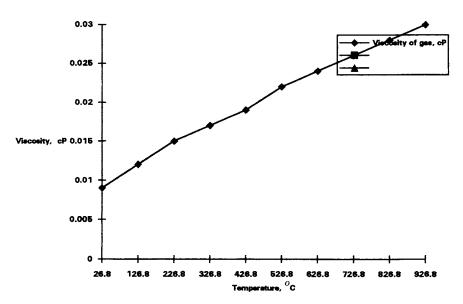


Figure 2-12. Viscosity of ethane as a function of temperature.

Liquid-Phase Diffusion Coefficients

Diffusion coefficients in liquid phases depend on concentration and are valid for dilute solutions, that is, solute concentrations no greater than 10%. Also, the lower the solute concentration, the more accurate the calculated coefficients. For a binary mixture of solute A in solvent B, the diffusion coefficient can be represented as D_{AB}^{0} for concentrations of A up to 5 or 10 mole percent [20]. A number of correlations have been proposed for predicting D_{AB}^{0} in dilute liquid solutions [21,22,23]. Here, the Wilke-Chang method is employed for estimating D_{AB}^{0} . This can be expressed as:

$$D_{AB}^{0} = \frac{7.4 \times 10^{-8} (\phi M_{wB})^{0.5} T}{\mu_{B} \bullet v_{A}^{0.6}}$$
(2-19)

where D_{AB}^{0} = Diffusion coefficient of solute A at very low concentrations in solvent B, cm²/s

- M_{wB} = molecular weight of solvent B
 - T = absolute temperature, K
 - $\mu_{\rm B}$ = viscosity of solvent B, cP
 - $v_A = molal volume of solute A at its normal boiling tempera$ ture, cm³/g-mol
 - ϕ = association factor of solvent B, dimensionless
 - = 2.6 for water (some workers recommend 2.26)
 - = 1.9 for methanol
 - = 1.5 for ethanol
 - = 1.0 for unassociated solvents

The Le Bas additive volumes given by Reid *et al.* [2] can be used for the value of V_A , if no experimental data are available.

Gas-Phase Diffusion Coefficients

Diffusion coefficients for non-polar gases may be estimated from Chapman and Cowling [24], Wilke and Lee [25] and Fuller, Schettler, and Giddings [26]. Here, the Fuller, Schettler, and Giddings's method can be expressed as:

$$D_{AB}^{0} = \frac{10^{-3} T^{1.75} \left\{ \frac{M_{wA} + M_{wB}}{M_{wA} \bullet M_{wB}} \right\}^{0.5}}{P \left\{ \left(\sum v \right)_{A}^{1/3} + \left(\sum v \right)_{B}^{1/3} \right\}^{2}}$$
(2-20)

where $D_{AB}^{0} = diffusion$ coefficient for a binary mixture of gases A and B, cm²/s $M_{wA}, M_{wB} =$ molecular weights of A and B $V_{A}, V_{B} =$ atomic diffusion volumes T = temperature, K P = pressure, atmospheres

The summation, $\sum v$, of the diffusion volume coefficients for components A and B is shown in Table 2-3. The computer program PROG23

· ·		·	
Atomic a	and Structural Diffu	sion-Volume Incre	ments, v
С	16.5	CI*	19.5
Н	1.98	S*	17.0
0	5.48	Aromatic ring	-20.2
N*	5.69	Heterocyclic ring	-20.2
Diff	usion Volumes for	Simple Molecules,	Σv
H ₂	7.07	СО	18.9
D ₂	6.70	CO ₂	26.9
Не	2.88	N ₂ O	35.9
N ₂	17.9	NH ₃	14.9
O ₂	16.6	H ₂ O	12.7
Air	20.1	CCl ₂ F [*] ₂	114.8
Ar	16.1	SF ₆ *	69.7
Kr	22.8	Cl [*]	37.7
Xe*	37.9	Br [*] ₂	67.2
		SO ₂ *	41.1

Table 2-3 Atomic Diffusion Volumes for Use in Calculating D_{AB} by the Correlation of Fuller-Schettler-Giddings

*Based on a few data points

calculates the liquid diffusion coefficient at varying molal volume of solute A, and the gas diffusion coefficient.

COMPRESSIBILITY Z-FACTOR OF NATURAL GASES

Many petroleum engineering and process design calculations dealing with natural gases require knowledge of deviation factors or compressibility Z factors. Experimental data from pressure-volume-temperature (P-V-T) measurements are seldom available. The Z-factors are available in charts [27] or tables as a function of pseudo-reduced temperatures T_r and pressures P_r . However, use of these charts is often time consuming and involves complex calculations.

Computer programs [28,29] for calculating the Z factors have been developed solely as a function of temperature and pressure of the gas. Furthermore, numerical methods and mathematical representations of the charts have been used to estimate the Z factors. Takacs [30] reviewed the various methods of estimating the Z-factors. Here, use is made of a modified form of the method developed by Awoseyin [31]. This method gives a compressibility factor to within 5% for natural hydrocarbon gases with specific gravities between 0.5 and 0.8 and for pressures up to 5000 psia. The Z factor can be expressed as:

$$Z = F_{1} \left\{ \frac{1}{\left[1 + \frac{\left(A_{6} \cdot P \cdot 10^{[1.785S_{g}]}\right)}{T^{3.825}}\right]} + F_{2} \cdot F_{3} \right\} + F_{4} + F_{5}$$
(2-21)

where

$$F_{I} = P(0.251S_{g} - 0.15) - 0.202S_{g} + 1.106$$
(2-22)

$$F_2 = 1.4e^{\{-0.0054(T-460)\}}$$
(2-23)

$$F_{3} = A_{1}P^{5} + A_{2}P^{4} + A_{3}P^{3} + A_{4}P^{2} + A_{5}P$$
(2-24)

$$F_{4} = \left\{ 0.154 - 0.152S_{g} \right\} P^{(3.18S_{g} - 1.0)} e^{(-0.5P)} - 0.02$$
 (2-25)

$$\mathbf{F}_{5} = 0.35 \left\{ (0.6 - \mathbf{S}_{g}) e^{[-1.039(P-1.8)^{2}]} \right\}$$
(2-26)

The values of the constants A_1 , A_2 , A_3 , A_4 , A_5 , and A_6 are:

 $A_{1} = 0.001946$ $A_{2} = -0.027635$ $A_{3} = 0.136315$ $A_{4} = -0.23849$ $A_{5} = 0.105168$ $A_{6} = 3.44 \times 10^{8}$

The specific gravity, S_g , of natural gas can be calculated from its density or molecular weight. This is expressed as the ratio of the gas density at 60°F and 1 atmosphere (14.7psia), $\rho_{gas,60^\circ F}$, to the density of air, $\rho_{air,60^\circ F}$, under the same conditions.

$$S_{g} = \frac{\text{density of gas}}{\text{density of air}}$$
$$= \frac{\rho_{\text{gas},60^{\circ}\text{F}}}{\rho_{\text{air},60^{\circ}\text{F}}}$$
(2-27)

Using the molecular weight of the gas, S_g , can be expressed as:

$$S_{g} = \frac{\text{molecular weight of gas}}{\text{molecular weight of air}}$$
$$= \frac{M_{w, \text{ gas}}}{M_{w, \text{ air}}}$$
(2-28)

Appendix A2 shows the computer outputs of the Z-factor as a function of temperature, pressure and specific gravity. The computer program PROG24 calculates the compressibility Z-factor of natural gases as a function of specific gravity and temperature. Figure 2-13 gives a plot of the compressibility Z-factor of natural gases at 60°F as a function of pressure and specific gravity. Tables 2-1 to 2-25 in Appendix B give results of Z-factors at varying specific gravity and temperature.

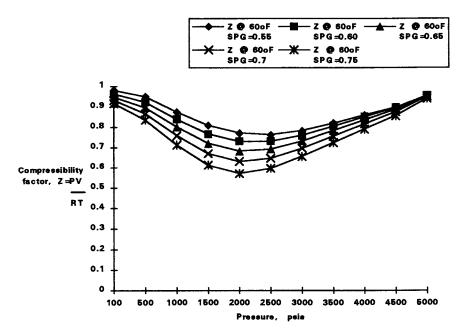


Figure 2-13. Compressibility factor of natural gases as a function of pressure and specific gravity.

PROBLEMS AND SOLUTIONS

Problem 2-1

Using the Wilke-Chang relation, determine the value of the infinitedilution diffusion coefficient of propane in chlorobenzene at 0°C. $M_B = 112.56$ (chlorobenzene), $\mu = 1.05$ cP, $V_A = 74.5$ (cm³/(g – mol)) (propane), T = 273.15K and $\phi = 1.0$.

Solution

The computer program PROG23 calculates the diffusion coefficient of propane in chlorobenzene at varying molal volume of solute A (propane). At $V_A = 74.5(\text{cm}^3/(\text{g}-\text{mol}))$, the computed value of the diffusion coefficient, $D_{AB}^0 = 1.54 \times 10^{-5} (\text{cm}^2/\text{s})$.

Problem 2-2

Calculate the diffusion coefficient of allyl chloride (AC) in air at 25°C and 1 atmosphere. An experimental value is 0.0975cm²/s.

Solution

Using the correlation of Fuller-Schettler-Giddings, values of atomic diffusion volumes of $\sum v$ from Table 2-3 are:

$$\sum v(AC) = 3(C) + 5(H) + Cl$$

= 3(16.5) + 5(19.8) + 19.5
= 78.9
$$\sum v(Air) = 20.1$$

Molecular weight of air, $M_{wA} = 29.0$ Molecular weight of allyl chloride, $M_{wB} = 76.53$

The computer program PROG23 determines the diffusion coefficient of the gas phase. The computed diffusion coefficient of allyl chloride is 0.0952cm²/s. Table 2-4 shows the computer output of diffusion coefficients for the liquid and gas phases.

Table 2-4Computer Output of Diffusion Coefficients for Liquid andGas Phases Diffusion Coefficients for Liquids and Gases

	IFFUSION COEFFICIENT OF L	IQUID PRASE
NUMBER	MOLAL VOLUME OF SOLUTE A cm^3/g-mol	DIFFUSION COEFFICIENT cm^2/s.
1	10.00	.5130E-04
2	20.00	.3385E-04
3	40.00	.2233E-04
4	60.00	.1751E-04
5	74.50	.1538E-04
6	80.00	.1473E-04
7	100.00	.1289E-04
8	200.00	.8502E-05
9	400.00	.5609E-05
10	600.00	.4398E-05 .3237E-05
11 12	1000.00 2000.00	.2136E-05
D:	IFFUSION COEFFICIENT OF G	as phase

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PROGRAM PROG21

с	THIS PROGRAM CALCULATES THE PHYSICAL PROPERTIES OF WATER
с с с с	AT A TEMPERATURE RANGE FROM OOC TO 5000C. THE CRITICAL TEMPERATURE=374.120C.
	THE PROGRAM WILL CALCULATE THE FOLLOWING PHYSICAL PROPERTIES 1. DENSITY DEN g/cm^3. 2. VISCOSITY VIS cP 3. HEAT CAPACITY HC cal/g.oC. 4. THERMAL CONDUCTIVITY THER (micro-cal /s.cm.K) 5. SURFACE TENSION ST dyne/cm. 6. VAPOR PRESSURE VP mm Hg. 7. HEATS OF VAPORIZATION HV cal/g
	DIMENSION T(1:50), DEN(1:50), VIS(1:50), HC(1:50) DIMENSION THER(1:50), ST(1:50), TK(1:50), VP(1:50), HV(1:50) INTEGER N
с	OPEN FILE TO PRINT THE RESULTS
	OPEN (UNIT = 1, FILE = 'PRN')
c c	INPUT THE NUMBER OF DATA AND THE VALUES OF THE TEMPERATURE FOR WHICH THE PHYSICAL PROPERTIES ARE CALCULATED.
	DATA N /10/
	DATA T(1)/0.0/, T(2)/25.0/, T(3)/50.0/, T(4)/100.0/ DATA T(5)/150.0/, T(6)/200.0/, T(7)/250.0/, T(8)/300.0/ DATA T(9)/350.0/, T(10)/374.2/
100 *	WRITE (1, 100) FORMAT (//, 1H , 25X, 'PHYSICAL PROPERTY DATA OF WATER', /1H ,78(1H*))
110	WRITE (1, 110) Format (//, 40X, 'temperature oC')
120	DO I = 1, N WRITE (1, 120) I, T(I) FORMAT (30X, I2, 10X, F8.2) END DO
125	WRITE (1, 125) Format (78(1H-))
с	CONVERT THE TEMPERATURE FROM DEGREE CENTIGRADE TO KELVIN.
	DO I = 1, N TK(I) = T(I) + 273.16 END DO

J = 1 GO TO (11, 22, 33, 44, 55, 66, 77), J 5 с CALL THE PROGRAM TO CALCULATE THE DENSITY OF WATER CALL PROG1 (T, DEN, N) 11 WRITE (1, 130) FORMAT (//, 1H , 25X, 'DENSITY OF WATER: 0oC - 374.2oC', /1H ,78(1H*)) 130 WRITE (1, 140) FORMAT (/,20X, ' TEMPERATURE oC', 10X, 'DENSITY g/cm.^3', 140 /, 78(1H-),/) DO I = 1, N WRITE (1, 150) I, T(I), DEN(I) FORMAT (10X, I2, 10X, F6.1, 10X, F12.3) 150 END DO WRITE (1, 155) FORMAT (78(1H-)) 155 J = J + 1IF (J.EQ. 2) THEN GO TO 5 ELSE GO TO 900 ENDIF с CALL THE PROGRAM TO CALCULATE WATER VISCOSITY CALL PROG2 (TK, VIS, N) 22 160 170 DO I = 1, N WRITE (1, 180) I, T(I), VIS(I) FORMAT (10X, I2, 10X, F6.1, 10X, F12.3) 180 END DO WRITE (1, 185) FORMAT (78(1H-)) 185 J = J + 1IF (J .EQ. 3) THEN GO TO 5 ELSE GO TO 900 ENDIF

с	CALL THE PROGRAM TO CALCULATE HEAT CAPACITY OF WATER
33	CALL PROG3 (TK, HC, N)
190 *	WRITE (1, 190) FORMAT (//, 1H , 25X, 'HEAT CAPACITY OF WATER: 0oC - 350oC', /1H ,78(1H*))
200 *	<pre>wRITE (1, 200) FORMAT (/, 20X,'TEMPERATURE oc',10X,'HEAT CAPACITY cal/gm.oC',</pre>
210	DO I = 1, N-1 WRITE (1,210) I, T(I), HC(I) FORMAT (10X, I2, 10X, F6.1, 10X, F12.3) END DO
215	WRITE (1, 215) Format (78(1H-))
	J = J + 1 IF (J .EQ. 4) THEN GO TO 5 ELSE GO TO 900 ENDIF
с	CALL THE PROGRAM TO CALCULATE THE THERMAL CONDUCTIVITY OF WATER
44	CALL PROG4 (TK, THER, N)
220 *	WRITE (1, 220) FORMAT (//, 25X, 'THERMAL CONDUCTIVITY OF WATER: 0oC - 350oC', /1H , 78(1H*))
230 240	WRITE (1, 230) FORMAT (/, 20X, 'TEMPERATURE oC', 5X,'THERMAL CONDUCTIVITY',\) WRITE (1, 240) FORMAT (' micro-cal/s.cm.K', /, 78(1H-), /)
250	DO I = 1, N-1 WRITE (1, 250) I, T(I), THER(I) FORMAT (10X, I2, 10X, F6.1, 10X, F12.3) END DO
255	WRITE (1, 255) Format (78(1H-))
	J = J + 1 IF (J .EQ. 5) THEN GO TO 5 ELSE GO TO 900 ENDIF
c	CALL THE PROGRAM TO CALCULATE THE SURFACE TENSION OF WATER (Page 219, Yaws et. al.)

C WATER (Page 219, Yaws et. al.)

55	CALL PROG5 (TK, ST, N)
260 *	<pre>WRITE (1, 260) FORMAT (//, 1H , 25X, 'SURFACE TENSION OF WATER: 0oC - 374.2oC',</pre>
270 *	<pre>wRITE (1, 270) FORMAT (/, 20X, 'TEMPERATURE oC',10X,'SURFACE TENSION:dyne/cm.',</pre>
280	DO I = 1, N WRITE (1, 280) I, T(I), ST(I) FORMAT (10X, I2, 10X, F6.1, 10X, F12.3) END DO
285	WRITE (1, 285) Format (78(1H-))
	J = J + 1 IF (J .EQ. 6) THEN GO TO 5 ELSE GO TO 900 ENDIF
c c	CALL THE PROGRAM TO CALCULATE THE VAPOR PRESSURE OF SATURATED LIQUIDS (PAGE 214 Yaws et. al.)
66 290 *	CALL PROG6 (TK, VP, N) WRITE (1, 290) FORMAT(//, 1H , 25X, 'VAPOR PRESSURE OF WATER: 0oC - 374.2 oC', /1H , 78(1H*))
300 *	WRITE (1, 300) FORMAT (/,20X,'TEMPERATURE oC', 10X,'VAPOR PRESSURE: mm Hg', /, 78(1H-), /)
310	DO I = 1, N WRITE (1, 310) I, T(I), VP(I) FORMAT (10X, I2, 10X, F6.1, 10X, F12.3) END DO
315	WRITE (1, 315) Format (78(1H-))
	J = J + 1 IF (J.EQ. 7) THEN
	GO TO 5 ELSE GO TO 900 ENDIF

77 320	CALL PROG7 (TK, HV, N) WRITE (1, 320) FORMAT(//, 1H ,25X,'HEAT-OF-VAPORIZATION OF WATER:00C-374.2 oC',
*	/1H , 78(1H*))
330 *	<pre>wRITE (1, 330) FORMAT(//,20X,'TEMPERATURE oC',10X,'HEAT-OF-VAPORIZATION:cal/g',</pre>
340	DO I = 1, N WRITE (1, 340) I, T(I), HV(I) FORMAT (10X, I2, 10X, F6.1, 10X, F12.3) END DO
345	WRITE (1, 345) Format (78(1H-))
	J = J + 1 IF (J .EQ. 8) THEN GO TO 900 ELSE ENDIF
900	CONTINUE
с	FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR (12)
	CLOSE (UNIT = 1) STOP END
с с с	**************************************
	SUBROUTINE PROG1 (T, DENL, N) DIMENSION T(1:50), DENL(1:50), TR(1:50), SUM(1:50)
	INTEGER N
c c	CRITICAL TEMPERATURE : TC=374.2 CALCULATE THE REDUCED TEMPERATURE : TR=T/TC
	TC = 374.2 DO I = 1, N TR(I) = T(I)/TC END DO
с	CORRELATION CONSTANTS OF WATER
<u> </u>	A = 0.3471 B = 0.2740
c c	CALCULATE THE DENSITY OF WATER
	DO 10 I = 1, N

ELSE SUM(I) = SUM(I) **0.2857DENL(I) = A*(B**(-SUM(I)))ENDIF 10 CONTINUE RETURN END C C ******** THIS PROGRAM CALCULATES THE VISCOSITY OF WATER FROM 0oC TO 374.2oC PAGE 213, YAWS et. al. С с *************** SUBROUTINE PROG2 (TK, VIS1, N) DIMENSION TK(1:50), VIS1(1:50), VIS2(1:50) INTEGER N с CORRELATION CONSTANTS OF WATER A = -10.73B = 1828.0C = .01966 D = -.00001466с CALCULATE THE VISCOSITY DO I = 1, N VIS2(I) = A + (B/TK(I)) + (C*TK(I)) + (D*TK(I)**2) VIS1(I) = 10**(VIS2(I))END DO RETURN END c c THIS PROGRAM CALCULATES THE HEAT CAPACITY OF WATER FROM OC TO 3500C (PAGE 221 Yaws et. al.) c c SUBROUTINE PROG3 (TK, HC1, N) DIMENSION TK (1:50), HC1 (1:50) CORRELATION CONSTANTS OF WATER с A = 0.6741B = .002825C = -.000008371D = .00000008601 с CALCULATE THE HEAT CAPACITY DO I = 1, N HCl(I) = A + (B*TK(I)) + (C*TK(I)**2) + (D*TK(I)**3)END DO RETURN

```
c
c
        THIS PROGRAM CALCULATES THE THERMAL CONDUCTIVITY OF WATER
        C
        SUBROUTINE PROG4 (TK, THERM1, N)
        DIMENSION TK(1:50), THERM1(1:50)
        INTEGER N
        CORRELATION CONSTANTS OF WATER
С
        A = -916.62
        B = 12.5473
        C = -.015212
C
C
        CALCULATE THE THERMAL CONDUCTIVITY OF WATER
        DO I = 1, N + 1
THERM1 (I) = A + (B*TK(I)) + (C*TK(I)**2)
        END DO
        RETURN
        END
        С
С
        THIS PROGRAM CALCULATES THE SURFACE TENSION OF WATER
c
c
        FROM 0oC TO 374.20C.
        SURFACE TENSION ST=ST1*((TC-T)/(TC-T1))^N1
        FOR 0 < T < 100oC, ST1=71.97, TC=374.20C, T1=25.00C.
FOR 10 < T < 374.20C, ST1=58.91, TC=374.20C, T1=100oC.
С
С
                                                               ......
C
        SUBROUTINE PROG5 (TK, ST, N)
DIMENSION TK(1:50), ST(1:50)
        REAL N1, N2
с
        CALCULATE THE SURFACE TENSION BETWEEN OOC AND 500C.
        ST1 = 71.97
        TC = 374.2 + 273.16
T1 = 25.0 + 273.16
        N1 = 0.8105
        DO I = 1, 3
ST(I) = ST1 * ((TC-TK(I))/(TC-T1))**N1
        END DO
с
        CALCULATE THE SURFACE TENSION BETWEEN 100oC AND 374.2oC.
        ST2 = 58.91
        T2 = 100.0 + 273.16
        N2 = 1.169
        DO I = 4, N
ST(I) = ST2 * ((TC-TK(I))/(TC-T2))**N2
        END DO
```

END

```
RETURN
END
```

```
С
С
С
С
       THIS PROGRAM CALCULATES THE VAPOR PRESSURE OF WATER FROM
       0oc To 374.2 oc.
с
       SUBROUTINE PROG6 (TK, VP1, N)
DIMENSION TK (1:50), VP1 (1:50)
c
c
       CORRELATION CONSTANTS OF WATER
       A = 16.373
       B = -2818.6
C = -1.6908
       D = -0.0057546
       E = 0.0000040073
c
c
       CALCULATE THE VAPOR PRESSURE
       DO I = 1, N
VP1(I) = A + (B/TK(I))+(C*ALOG10(TK(I)))+(D*TK(I))+(E*TK(I)**2)
VP1(I) = 10**VP1(I)
       END DO
       RETURN
       END
       с
с
       THIS PROGRAM CALCULATES THE HEAT OF VAPORIZATION OF WATER
c
c
       FROM 0oC TO 374.2 oC.
       *****
       SUBROUTINE PROG7 (TK, HV1, N)
       DIMENSION TK(1:50), HV1(1:50)
с
       CORRELATION CONSTANTS OF WATER
       THV = 538.7
       T1 = 100.0 + 273.16
TC = 374.2 + 273.16
       R = 0.38
¢
       CALCULATE THE HEAT OF VAPORIZATION
      DO I = 1, N
HV1(I) = THV*(((TC-TK(I))/(TC-T1))**R)
       END DO
       RETURN
       END
```

PROGRAM PROG22

C	DIMENSION T (1:50), HF (1:50), GF (1:50), HC (1:50) DIMENSION THER (1:50), VIS (1:50), TK (1:50)
	INTEGER N
с	OPEN FILE TO PRINT THE RESULTS.
	OPEN (UNIT = 1, FILE = 'PRN')
с с	INPUT THE NUMBER OF DATA AND THE VALUES OF THE TEMPERATURE FOR WHICH THE PHYSICAL PROPERTIES ARE CALCULATED.
	DATA N /13/
	DATA TK(1)/300.0/, TK(2)/400.0/, TK(3)/500.0/, TK(4)/600.0/ DATA TK(5)/700.0/, TK(6)/800.0/, TK(7)/900.0/, TK(8)/1000.0/ DATA TK(9)/1100.0/, TK(10)/1200.0/, TK(11)/1300.0/ DATA TK(12)/1400.0/, TK(13)/1500.0/
100 *	WRITE (1, 100) FORMAT (//, 1H , 25X, 'PHYSICAL PROPERTY DATA OF GASES', /1H ,78(1H*))
110	WRITE (1, 110) FORMAT (//, 40X, 'TEMPERATURE RANGE (300 - 1500K)')
115	WRITE (1, 115) Format (78(1H-))
120	DO I = 1, N WRITE (1, 120) I, TK(I) FORMAT (30X, I2, 10X, F8.2) END DO
125	WRITE (1, 125) Format (78(1H-))
	J = 1
5	GO TO (11, 22, 33, 44, 55), J

с	CALL THE PROGRAM TO CALCULATE THE HEAT OF FORMATION OF A GAS
11	CALL PROG1 (TK, HF, N)
130 *	WRITE (1, 130) FORMAT (//, 1H , 25X, 'HEAT OF FORMATION OF METHANE', /1H , 78(1H*), /)
140 *	<pre>WRITE (1, 140) FORMAT (20X,' TEMPERATURE K', 5X,'HEAT OF FORMATION kcal/mol',</pre>
150	DO I = 1, N WRITE (1, 150) I, TK(I), HF(I) FORMAT (10X, I2, 10X, F6.1, 10X, F12.3) END DO
155	WRITE (1, 155) Format (78(1H-))
	J = J + 1 IF (J .EQ. 2) THEN GO TO 5 ELSE GO TO 900 ENDIF
c c	CALL THE PROGRAM TO CALCULATE THE FREE ENERGY OF FORMATION OF A GAS
22	CALL PROG2 (TK, GF, N)
160 165	WRITE (1, 160) FORMAT (//, 1H , 25X, 'THE FREE ENERGY OF FORMATION OF',\) WRITE (1, 165) FORMAT(' ETHYLENE kcal/mol.', /1H ,78(1H*), /)
170 F	WRITE (1, 170) ORMAT(20X,'TEMPERATURE K',5X,'FREE ENERGY OF FORMATION:kcal/mol')
175	WRITE (1, 175) Format (78(1H-), /)
180	DO I = 1, N WRITE (1, 180) I, TK(I), GF(I) FORMAT (10X, I2, 10X, F6.1, 10X, F12.3) END DO
185	WRITE (1, 185) Format (78(1H-))
	J = J + 1 IF (J .EQ. 3) THEN GO TO 5 ELSE GO TO 900 ENDIF
с	CALL THE PROGRAM TO CALCULATE HEAT CAPACITY OF CARBON DIOXIDE

33 190 *	CALL PROG3 (TK, HC, N) WRITE (1, 190) FORMAT (//, 1H , 25X, 'HEAT CAPACITY OF CARBON DIOXIDE', /1H ,78(1H*))
200 *	<pre>WRITE (1, 200) FORMAT (20%,'TEMPERATURE K', 5%,'HEAT CAPACITY cal/gm-mol.K',</pre>
210	DO I = 1, N WRITE (1, 210) I, TK(I), HC(I) FORMAT (10X, I2, 15X, F6.1, 5X, F12.3) END DO
215	WRITE (1, 215) Format (78(1H-))
	J = J + 1 IF (J .EQ. 4) THEN GO TO 5 ELSE GO TO 900 ENDIF
с	CALL THE PROGRAM TO CALCULATE THE THERMAL CONDUCTIVITY OF GASES
44	CALL PROG4 (TK, THER, N)
220 *	WRITE (1, 220) FORMAT(//,25X,'THERMAL CONDUCTIVITY OF PROPANE:26.8oC-1026.8oC', /1H , 78(1H*))
230 235	<pre>WRITE (1, 230) FORMAT (20X,'TEMPERATURE oC', 5X,'THERMAL CONDUCTIVITY',\) WRITE (1, 235) FORMAT(' micro-cal/s.cm.K', /, 78(1H-), /)</pre>
	DO I = 1, $N-2$
240	T(I) = TK(I) - 273.16 WRITE (1, 240) I, T(I), THER(I) FORMAT (10X, I2, 10X, F6.1, 10X, F12.3) END DO
245	WRITE (1, 245) Format (78(1H-))
	J = J + 1 IF (J .EQ. 5) THEN GO TO 5 ELSE GO TO 900 ENDIF
с	CALL THE PROGRAM TO CALCULATE THE VISCOSITY OF ETHANE

```
55
        CALL PROG5 (TK, VIS, N)
       250
       260
        DO I = 1, N - 2
T(I) = TK(I) - 273.16
WRITE (1, 270) I, T(I), VIS(I)
FORMAT (10X, I2, 10X, F6.1, 10X, F12.3)
 270
        END DO
       WRITE (1, 275)
FORMAT ( 78(1H-))
 275
        \mathbf{J}=\mathbf{J}+\mathbf{1}
        IF (J .EQ. 6) THEN
        GO TO 900
        ELSE
        ENDIF
 900
        CONTINUE
с
        FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
        WRITE (1, *) CHAR (12)
        CLOSE (UNIT = 1)
        STOP
        END
        С
        THIS PROGRAM CALCULATES THE HEAT OF FORMATION OF METHANE
"PHYSICAL PROPERTIES BY C.L. YAWS et. al. PAGE 200
с
c
c
        SUBROUTINE PROG1 (TK, HF1, N)
DIMENSION TK(1:50), HF1(1:50)
        INTEGER N
с
        CORRELATION CONSTANTS OF METHANE
        A = -15.4
        B = -0.00959
        C = 0.0000035
с
        CALCULATE THE HEAT OF FORMATION OF METHANE, kcal/mol.
        DO I = 1, N
        HF1(I) = A + (B*TK(I)) + (C*TK(I)**2)
END DO
        RETURN
        END
```

```
С
       С
       THIS PROGRAM CALCULATES THE FREE ENERGY OF FORMATION OF
       ETHYLENE PAGE 202, YAWS et. al.
с
ē
                                   *****
       SUBROUTINE PROG2 (TK, GF1, N)
DIMENSION TK(1:50), GF1(1:50)
       INTEGER N
С
       CORRELATION CONSTANTS OF ETHYLENE
       A = 10.4
       B = 0.0181
С
       CALCULATE THE FREE ENERGY OF FORMATION
       DO I = 1, N
GF1(I) = A + (B*TK(I))
       END DO
       RETURN
       END
с
       THIS PROGRAM CALCULATES THE HEAT CAPACITY OF CARBON DIOXIDE
0oC-350oC (PAGE 198 Yaws et. al.)
C
C
C
       SUBROUTINE PROG3 (TK, HC1, N)
       DIMENSION TK(1:50), HC1(1:50)
с
       CORRELATION CONSTANTS OF CARBON DIOXIDE
       A = 5.41
       B = 0.0154
       C = -0.00000994
       D = 0.0000000242
с
       CALCULATE THE HEAT CAPACITY
       DO I = 1, N
HCl(I) = A + (B*TK(I))+(C*TK(I)**2)+(D*TK(I)**3)
       END DO
       RETURN
       END
с
       с
       THIS PROGRAM CALCULATES THE THERMAL CONDUCTIVITY OF PROPANE
С
       PAGE 208 Yaws et al.
с
       SUBROUTINE PROG4 (TK, THERM1, N)
DIMENSION TK(1:50), THERM1(1:50)
       INTEGER N
С
       CORRELATION CONSTANTS OF PROPANE
```

```
A = 4.438
B = -0.01122
C = 0.0005198
        D = -0.000002008
с
        CALCULATE THE THERMAL CONDUCTIVITY OF PROPANE
       DO I = 1, N - 2
THERM1(I) = A + (B*TK(I))+(C*TK(I)**2)+(D*TK(I)**3)
        END DO
        RETURN
        END
c
c
        THIS PROGRAM CALCULATES THE VISCOSITY OF ETHANE FROM
        0 - 1000 oC. PAGE 210 Yaws et al.
C
C
       SUBROUTINE PROG5 (TK, VIS1, N)
DIMENSION TK(1:50), VIS1(1:50)
       CORRELATION CONSTANTS OF ETHANE
с
        A = 5.576
       B = 0.3064
       C = -0.00005307
       CALCULATE THE VISCOSITY
С
       DO I = 1, N-2
VIS1(I) = (A + (B*TK(I))+(C*TK(I)**2))*0.0001
        END DO
        RETURN
        END
```

PROGRAM PROG23

с с с	**************************************		
с с	DIFFUSION COEFFICIENT FOR LIQUID PHASE		
с с с	THE PROGRAM USES THE WILKE-CHANG METHOD TO CALCULATE THE DIFFUSION COEFFICIENT OF A BINARY MIXTURE OF SOLUTE A IN SOLVENT B.		
00000000	DABL = DIFFUSION COEFFICIENT OF LIQUID PHASE cm^2/s. MBL = MOLECULAR WEIGHT OF SOLVENT B. TKL = ABSOLUTE TEMPERATURE, K VISBL = VISCOSITY OF SOLVENT B, CP VA = MOLAL VOLUME OF SOLUTE A AT ITS NORMAL BOILING TEMPERATURE, cm^3/g-mol. FACTOR = ASSOCIATION FACTOR OF SOLVENT B, DIMENSIONLESS		
с	DIFFUSION COEFFICIENT FOR GAS PHASE		
с	THE PROGRAM USES THE FULLER, SCHETTLER AND GIDDLINGS METHOD		
с с с с с с с с с	DABG = DIFFUSION COEFFICIENT OF GAS PHASE cm^2/s. TKG = TEMPERATURE, KELVIN MWA = MOLECULAR WEIGHT OF A. MWB = MOLECULAR WEIGHT OF B VAG = ATOMIC DIFFUSION VOLUME OF A VBG = ATOMIC DIFFUSION VOLUME OF B PRES = PRESSURE, ATMOSPHERE ***********************************		
с	OPEN FILE TO PRINT THE RESULTS. OPEN (UNIT = 1, FILE = 'PRN')		
с	DATA FOR LIQUID PHASE DIFFUSION COEFFICIENT		
	DATA VA(1)/10.0/, VA(2)/20.0/, VA(3)/40.0/, VA(4)/60.0/ DATA VA(5)/74.5/, VA(6)/80.0/, VA(7)/100.0/, VA(8)/200.0/ DATA VA(9)/400.0/, VA(10)/600.0/, VA(11)/1000.0/, VA(12)/2000.0/ DATA TKL/273.15/, MBL/112.56/, VISBL/1.05/, FACTOR/1.0/		
с	DATA FOR GAS PHASE DIFFUSION COEFFICIENT DATA TKG/298.15/, MWA/29.0/, MWB/76.53/, VAG/20.1/, VBG/78.9/ DATA PRES/1.0/ WRITE (1, 100)		

145

100 J = 1 5 GO TO (11, 22), J с CALL THE PROGRAM TO CALCULATE THE DIFFUSION COEFFICIENT OF С LIQUID PHASE 11 110 CALL PROG1 (DABL, TKL, MBL, VISBL, VA, FACTOR, Y) 140 DO I = 1, 12 WRITE (1, 150) I, VA(I), DABL(I) FORMAT (10X, I2, 10X, F10.2, 15X, E12.4) 150 END DO WRITE (1, 160) FORMAT (78(1H-)) 160 J = J + 1IF (J .EQ. 2) THEN GO TO 5 ELSE GO TO 900 ENDIF C C CALL THE PROGRAM TO CALCULATE THE DIFFUSION COEFFICIENT OF A GAS PHASE 22 10 * CALL PROG2 (DABG, TKG, MWA, MWB, VAG, VBG, PRES) WRITE (1, 170) DABG 170 FORMAT (//, 15X, 'DIFFUSION COEFFICIENT:', E12.6, 3X, 'Cm^2/s.') J = J + 1IF (J .EQ. 2) THEN GO TO 900

```
ELSE
        ENDIF
 900
        CONTINUE
        WRITE (1, 180)
FORMAT (78(1H-))
 180
с
        FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
        WRITE (1, *) CHAR(12)
        CLOSE (UNIT = 1)
        STOP
        END
С
        THIS PROGRAM CALCULATES THE DIFFUSION COEFFICIENT OF THE LIQUID PHASE USING THE WILKE-CHANG METHOD.
С
c
c
                                                     ******
        SUBROUTINE PROG1 (DABL, TK, MB, VISB, VA, FACTOR, Y)
        DIMENSION VA(1:50), DABL(1:50), Y(1:50)
        REAL MB
с
        DO I = 1, 12
        SUM1 = (7.4*(FACTOR*MB)**0.5*TK)
SUM2 = (VISB*VA(I)**0.6)
DABL(I) = SUM1/SUM2
DABL(I) = DABL(I)/(10**8)
        Y(I) = TK/(DABL(I)*VISB*10**7)
        END DO
        RETURN
        END
С
        с
        THIS PROGRAM CALCULATES THE DIFFUSION COEFFICIENT OF THE
c
        GAS PHASE USING THE FULLER, SCHETTLER AND GIDDINGS METHOD
        SUBROUTINE PROG2 (DABG, TKG, MAG, MBG, VAG, VBG, PRES)
        REAL MAG, MBG
        SUM3 = ((MAG+MBG)/(MAG*MBG))**0.5
        SUM4 = (VAG**0.333+VBG**0.333)**2
        DABG = (TKG**1.75*0.001*SUM3)/(PRES*SUM4)
        RETURN
        END
```

PROGRAM PROG24

C C C	THIS PROGRAM CALCULATES THE COMPRESSIBILITY FACTOR OF NATURAL GAS AT A KNOWN SPECIFIC GRAVITY AND TEMPERATURE
с с с	**************************************
с с с	TF = GAS TEMPERATURE, oF. Z = GAS COMPRESSIBILITY FACTOR.
с	OPEN FILE TO PRINT THE RESULTS.
	OPEN (UNIT=1, FILE='PRN')
	DATA TF /60.0/, SPG /0.550/
9	WRITE (1, 100)
100 *	<pre>FORMAT (20X,'COMPRESSIBILITY FACTOR OF NATURAL GAS',</pre>
105	WRITE (1, 105) FORMAT (10X, 'TEMPERATURE, oF', 10X, 'SPECIFIC GRAVITY')
110	WRITE (1,110) TF, SPG FORMAT (10X, F6.0, 20X, F6.3,/)
	$\begin{array}{l} \lambda 1 = 0.001946 \\ \lambda 2 = -0.027635 \\ \lambda 3 = 0.136315 \\ \lambda 4 = -0.238489 \\ \lambda 5 = 0.105168 \\ \lambda 6 = 3.444*(10**8) \\ T = TF + 460.0 \end{array}$
120 *	<pre>WRITE (1, 120) FORMAT (10X,'PRESSURE psia.',6X,'COMPRESSIBILITY FACTOR Z',</pre>
	PP = 100.0
10	P1 = PP/1000.0
с	WRITE THE EXPRESSIONS FOR F1, F2, F3, F4, F5
	$\begin{array}{l} F1 &= P1 \star (0.251 \star SPG-0.150) - 0.202 \star SPG + 1.106 \\ F2 &= 1.4 \star EXP(-0.0054 \star TF) \\ F3 &= (A1 \star P1 \star \star 5) + (A2 \star P1 \star \star 4) + (A3 \star P1 \star \star 3) + (A4 \star P1 \star \star 2) + A5 \star P1 \\ VAL1 &= P1 \star \star (3.18 \star SPG-1.0) \\ VAL2 &= EXP(-0.5 \star P1) \\ F4 &= (VAL1 \star VAL2) \star (0.154 - 0.152 \star SPG) - 0.02 \\ VAL3 &= -1.039 \star ((P1 - 1.8) \star \star 2) \\ F5 &= 0.35 \star (0.6 - SPG) \star EXP(VAL3) \\ VAL4 &= 1 + (A6 \star P1 \star 10 \star (1.785 \star SPG))/(T \star \star 3.825) \\ Z &= F1 \star (1/VAL4 + (F2 \star F3)) + F4 + F5 \end{array}$

130	WRITE (1, 130) PP, Z Format (5X, F12.0, 15X, F12.5)
	PP = PP + 100.0
	IF (PP .GT. 5000.0) THEN GO TO 60 ELSE GO TO 10 ENDIF
с	PRINT THE RESULTS ON A NEW PAGE.
60 140	WRITE (1, 140) FORMAT (78(1H-))
150	WRITE (1, 150) FORMAT ('1')
	TF = TF + 10.0
	IF (TF .GT. 100.0) THEN GO TO 12 ELSE GO TO 9 ENDIF
12	TF = TF - 50.0 SPG = SPG + 0.05
	IF (SPG .GT. 0.8) THEN GO TO 70 ELSE GO TO 9 ENDIF
с	FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR(12)
70	CLOSE (UNIT = 1) STOP END

CHAPTER 3

Fluid Flow

INTRODUCTION

Transportation of fluids is important in the design of chemical process plants. In the chemical process industries (CPI), pipework and its accessories such as fittings, makeup 20% to 30% of the total design costs and 10% to 20% of the total plant investment. Maintenance requirements and energy usage in the shape of pressure drop (ΔP) in the fluids being pumped add to the cost. Also, these items escalate each year in line with inflation. As a result, sound pipe sizing practices can have a substantial influence on overall plant economics. It is the designer's responsibility to optimize the pressure drops in piping and equipment and to assess the most economical conditions of operations. Figures 3-1A and 3-1B illustrate piping layouts in a chemical plant.

The characteristics and complexity of flow pattern are such that most flows are described by a set of empirical or semi-empirical equations. These relate the pressure drop in the flow system as a function of flow rate, pipe geometry, and physical properties of the fluids. The aim in the design of fluid flow is to choose a line size and piping arrangement that achieve minimum capital and pumping costs. In addition, constraints on pressure drop and maximum allowable velocity in the process pipe should be maintained. These objectives require many trial and error computations, which can be performed well by a computer.

Flow of Fluids in Pipes

Pressure drop or head loss in a piping system is caused by fluid rising in elevation, friction, shaftwork (e.g., from a turbine) and turbulence due to sudden changes in direction or cross-sectional area. Figure 3-2

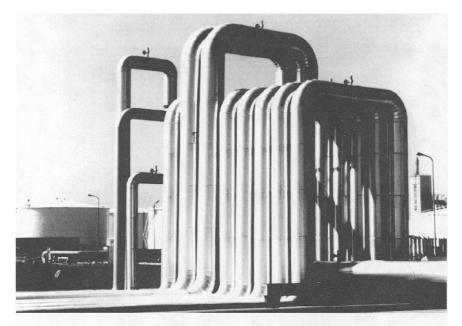


Figure 3-1A. Chemical plant piping layout.

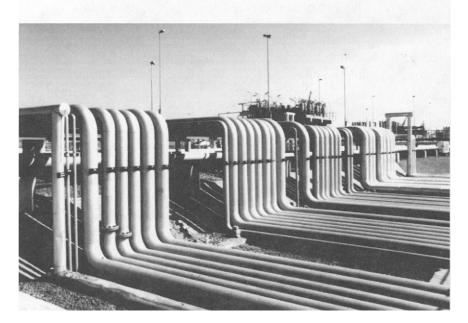


Figure 3-1B. Chemical plant piping layout. Source: I Chem E safer piping training package (courtesy of the I. Chem E., U.K.)

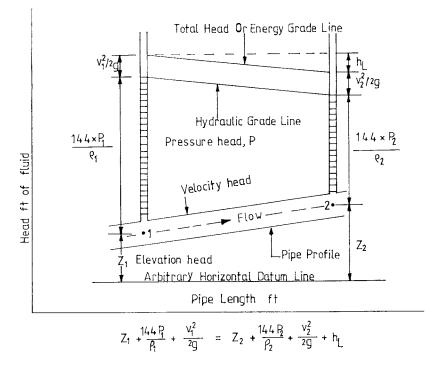


Figure 3-2. Distribution of fluid energy in a pipeline.

shows the distribution of energy between two points in a pipeline. Bernoulli's equation expresses the conservation of the sum of pressure, kinetic and potential energies.

$$\frac{1}{\rho} \int_{1}^{2} dP + \frac{1}{g_{c}} \int_{1}^{2} v dv + \int_{1}^{2} dz = 0$$
(3-1)

Integrating Equation 3-1 gives

$$\frac{1}{\rho}(\mathbf{P}_2 - \mathbf{P}_1) + \frac{\mathbf{v}_2^2 - \mathbf{v}_1^2}{2g_c} + (z_2 - z_1) = 0$$
(3-2)

where $P = \text{pressure of fluid, } lb_f/ft^2$ $\rho = \text{density of fluid, } lb_m/ft^3$ V = velocity of fluid, ft/s $g_c = \text{dimensional constant, } 32.174(lb_m/lb_f)(ft/s^2)$ Z = elevation of fluid, ftsubscript 1 = condition at initial point subscript 2 = condition at final point The first, second, and third terms in Equation 3-2 represent pressure head, velocity head, and static differences respectively. Equation 3-2 is used for investigating energy distributions or determining pressure differentials between any two points in a pipeline. Incorporating the head loss due to friction, h_L , with constant pipe diameter, i.e., $V_1 = V_2$, Equation 3-2 becomes

$$\frac{1}{\rho}(P_2 - P_1) + (Z_2 - Z_1) = h_L$$
(3-3)

Equation 3-3 shows that the head loss, h_L , is generated at the expense of pressure head or static head difference. The static head difference can be either negative or positive. However, for a negative static head difference,

$$\Delta P > h_{L} + (Z_{2} - Z_{1})$$
(3-4)

In general, pressure loss due to flow is the same whether the pipe is horizontal, vertical, or inclined. The change in pressure due to the difference in head must be considered in the pressure drop calculation.

EQUIVALENT LENGTH OF VARIOUS FITTINGS AND VALVES

The effects of bends and fittings, such as elbow, valves, tees, and reduction or enlargement of pipes are determined empirically through a fictituous equivalent length of straight pipe having the same diameter and they would develop the same pressure drop. The equivalent pipe length is the most convenient method for determining the overall ΔP in a pipe. The drawback to this approach is that, for a given fitting the equivalent length is not constant but depends on the Reynolds number, the pipe roughness, the pipe size, and the geometry of the fitting. The equivalent length is added to the length of actual straight pipe to give the total length of pipe.

$$L_{\text{Total}} = L_{\text{st}} + L_{\text{eq}} \tag{3-5}$$

where $L_{Total} = total length of pipe, ft$ $L_{st} = length of straight pipe, ft$ $L_{eg} = equivalent length of pipe, ft$

The pressure drop equation can be expressed as:

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$$\Delta \mathbf{P} = \left(\mathbf{f}_{\mathrm{D}} \frac{\mathbf{L}}{\mathbf{D}} + \sum \mathbf{K} \right) \rho \frac{\mathbf{v}^{2}}{2g_{\mathrm{c}}}$$
(3-6)

and the head loss equation can be expressed as:

$$\Delta H = \left(f_{\rm D} \frac{L}{D} + \sum K \right) \frac{v^2}{2g_{\rm c}}$$
(3-7)

where D = internal diameter of pipe, ft

 $f_D = Darcy friction factor$

K = Excess head loss

L = pipe length, ft

Excess Head Loss

K is a dimensionless factor defined as the excess head loss in a pipe fitting, and expressed in velocity heads. The velocity head is the amount of kinetic energy contained in a stream or the amount of potential energy required to accelerate a fluid to its flowing velocity. Most published K values apply to fully developed turbulent flow because at high Reynolds number, K is found to be independent of Reynolds number. However, the two-K technique includes a correction factor for low Reynolds number. Hooper [1] gives a detailed analysis of his method compared to others [2,3], and has shown that the two-K method is the most suitable for any pipe size. In general, the two-K method is independent of the roughness of the fittings, but it is a function of Reynolds number and of the exact geometry of the fitting. The method can be expressed as:

$$K = \frac{K_{\perp}}{N_{Re}} + K_{\infty} \left(1 + \frac{1}{d} \right)$$
(3-8)

where $K_1 = K$ for fitting at $N_{Re} = 1$

 $K_{\infty} = K$ for a large fitting at $N_{Re} = \infty$ d = internal diameter of attached pipe, inch

The conversion between equivalent pipe length and the resistance coefficient, K, can be expressed as:

$$K = f_{D} \frac{L_{eq}}{D}$$
(3-9)

Table 3-1 lists values of K_1 and K_{∞} for the two-K method.

Table 3-1Velocity Head Factors of Pipe Fittings

		Fi	tting Type	K ₁	K∞
		Standard (R/D = 1), screwed Standard (R/D = 1), flanged/welded Long-radius (R/D = 1.5, all types		800 800 800	0.40 0.25 0.20
Elbows	90°	Mitered elbows (R/D = 1.5)	1 Weld (90° angle) 2 Weld (45° angles) 3 Weld (30° angles) 4 Weld (22 1/2° angles) 5 Weld (18° angles)	1000 800 800 800 800 800	1.15 0.35 0.30 0.27 0.25
	45°	Standard (R/ D = 1), all types Long-radius (R/D = 1.5), all types Mitered, 1 weld, 45° angle Mitered, 2 weld, 22 1/2° angles		500 500 500 500	0.20 0.15 0.25 0.15
	180°	Standard ($R/D = 1$), screwed Standard ($R/D = 1$), flanged/welded Long radius ($R/D = 1.5$) all types		1000 1000 1000	0.60 0.35 0.30
Tees	Used as elbow	Standard, screwed Long-radius, screwed Standard, flanged or welded Stub-in-type branch		500 800 800 1000	0.70 0.40 0.80 1.00
	Run- through tee	Screwed Flanged or welded Stub-in-type branch		200 150 100	0.10 0.05 0.00
	Gate, ball, plug	Full line size, β Reduced trim, Reduced trim,	$\beta = 0.9$	300 500 1000	0.10 0.15 0.25
Valves	Globe, a	, standard , angle or Y-type ragm, dam type fly		1500 1000 1000 800	4.00 2.00 2.00 0.25
	Check	Lift Swing Tilting-disk		2000 1500 1000	10.00 1.50 0.50
			s for $R/D = 5$ pipe bends, flow through crosses.	45° to 18	80°. Us

Source: Hooper [1]

Pipe Reduction and Enlargement

The velocity head is $v^2/2g_c$. For any velocity profile the true velocity represents the integral of the local velocity head across the pipe diameter. It is found by dividing the volumetric flow rate by the cross-sectional area of the pipe and multiplying by a correction factor. For laminar flow this factor is 2; for turbulent flow the factor depends on both the Reynolds number and the pipe roughness and $(1 + 0.8f_D)$ [4]. Therefore, when the pipe size changes, the velocity head also changes. Because the velocity is inversely proportional to the flow area and thus to the diameter squared, K is inversely proportional to the velocity squared.

$$K_2 = K_1 \left(\frac{D_2}{D_1}\right)^4 \tag{3-10}$$

Although the potential energy provides the flowing fluid with kinetic energy at the pipe entrance, the kinetic energy is later recovered. This indicates that the measured pipe pressure will be lower than the calculated pressure by one velocity head. If the kinetic energy is not recovered at the pipe exit, the exit counts as a loss of one velocity head. Table 3-2 shows how K varies with changes in pipe size.

PRESSURE DROP CALCULATIONS FOR SINGLE-PHASE INCOMPRESSIBLE FLUIDS

The pressure drop of flowing fluids can be calculated from the Darcy friction factor:

$$\frac{\Delta P}{L} = \frac{f_{D}}{D} \bullet \frac{\rho V^{2}}{2g_{c}}$$
(3-11)
where f_{D} = Darcy friction factor
 ρ = density of fluid, lb_{m}/ft^{3}
 V = velocity of fluid in the pipe, ft/s
 g_{c} = dimensional constant, 32.174(lb_{m}/lb_{f})(ft/s²)
 L = length of pipe, ft

Friction Factor

The friction factor is related to the Reynolds number by a set of correlations and depends on whether the flow regime is laminar, transitional, or turbulent. The Reynolds number is:

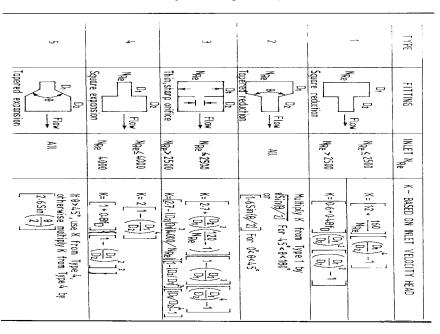


 Table 3-2

 Excess Head Loss 'K' Correlation for Fitting

 Caused by a Change in Pipe Size

$$N_{Re} = \frac{\rho VD}{\mu} = 50.6 \frac{Q\rho}{d\mu} = 5.31 \frac{W}{d\mu}$$
(3-12)

where Q = volumetric flowrate, gals/min W = Mass flow rate, lb/h μ = fluid viscosity, cP

For laminar flow with $N_{Re} \leq 2000$

$$f_{\rm D} = \frac{64}{N_{\rm Re}} \tag{3-13}$$

The Darcy friction factor is four times the Fanning friction factor, f_F , i.e., $f_D = 4f_F$. For fully developed turbulent flow regime in smooth and rough pipes, the Colebrook [5] equation or the Chen [6] equation can be used.

The Colebrook equation is expressed as:

$$\frac{1}{\sqrt{f_{\rm D}}} = -0.8686 \ln \left\{ \frac{\varepsilon}{3.7\rm{D}} + \frac{2.51}{\rm{N}_{\rm Re}\sqrt{f_{\rm D}}} \right\}$$
(3-14)

Equation 3-14 is implicit in f_D , as it cannot be rearranged to derive f_D directly and thus requires an iterative solution. Here, the Chen equation is used for calculating f_D , (i.e, $f_D = 4f_C$). The Chen equation is explicit and easier to use than Equation 3-14). This can be expressed as:

$$\frac{1}{\sqrt{f_c}} = -4 \log\left\{\frac{\varepsilon}{3.7D} - \frac{5.02}{N_{Re}}\log A\right\}$$
(3-15)

where

$$A = \frac{\varepsilon/D}{3.7} + \left(\frac{6.7}{N_{Re}}\right)^{0.9}$$

and

 ε = pipe roughness, ft

A detailed review of other explicit equations is given by Gregory and Fogarasi [7]. Different piping materials are often used in the chemical process industries, and at a high Reynolds number, the friction factor is affected by the roughness of the surface. This is measured as the ratio ϵ/D of projections on the surface to the diameter of the pipe. Glass and plastic pipe essentially have $\epsilon = 0$. Values of ϵ are shown in Table 3-3.

The Overall Pressure Drop

Designers involved in sizing process piping often apply trial and error procedure. The designer first selects a pipe size and then calculates the

Pipe Material	ε, ft
Riveted steel	0.003-0.03
Concrete	0.001-0.01
Wood stave	0.0006-0.003
Cast iron	0.00085
Galvanized iron	0.0005
Asphalted cast iron	0.0005
Commercial steel or wrought iron	0.00015
Drawn tubing	0.000005

Table 3-3 Values of Absolute Pipe Roughness

Reynolds number, friction factor, and coefficient of resistance. The pressure drop per 100 feet of pipe is then computed. For a given volumetric rate and physical properties of a single-phase fluid, ΔP_{100} for laminar and turbulent flows is:

laminar flow

$$\Delta P_{100} = 0.0273 \frac{\mu Q}{d^4}, \quad \text{psi / 100ft}$$
 (3-16)

turbulent flow

$$\Delta P_{100} = 0.0216 f_{\rm D} \frac{\rho Q^2}{d^5}, \quad \text{psi / 100 ft}$$
(3-17)

Alternatively, for a given mass flow rate and physical properties of a single-phase fluid, ΔP_{100} for laminar and turbulent flows respectively is:

laminar flow

$$\Delta P_{100} = 0.0034 \frac{\mu W}{d^4 \rho}, \quad psi / 100 ft$$
 (3-18)

turbulent flow

$$\Delta P_{100} = 0.000336 f_{\rm D} \frac{W^2}{d^5 \rho}, \quad \text{psi} / 100 \,\text{ft}$$
 (3-19)

Multiplying Equations 3-16, 3-17, 3-18, and 3-19 by the total length between two points and adding the pipe elevation yields the overall pressure drop.

$$\Delta P = \Delta P \bullet \frac{L_{\text{Total}}}{100} + \frac{\rho \Delta Z}{144}, \quad \text{psi}$$
(3-20)

Equation 3-20 is valid for compressible isothermal fluids of shorter lines where pressure drops are no more than 10% of the upstream pressure. In general, pipe size for a given flow rate is often selected on the assumption that the overall pressure drop is close to or less than the available pressure difference between two points in the line.

Nomenclature

A = pipe cross-sectional area, ft^2 d = internal pipe diameter, inch D = internal pipe diameter, ft

- f_c = Chen friction factor
- $f_p = Darcy friction factor$
- $g_c = dimensional constant 32.174 (lb_m/lb_f \cdot ft/s^2)$
- $h_L = head loss, ft$
- K = excess head loss for a fitting, velocity heads
- $K_1 = K$ for fitting at $N_{Re} = 1$, velocity heads
- $K_{\infty} = K$ for very large fitting at $N_{Re} = \infty$, velocity heads
- L_{eq} = equivalent length of pipe, ft
- L_{st} = actual length of pipe, ft
- L_{Total} = total length of pipe, ft
 - n = number of fittings
- N_{Re} = Reynolds number

$$\Delta P_{100}$$
 = pressure drop per 100 ft of pipe, psi/100 ft

- ΔP = overall pressure drop of pipe, psi
 - $\rho =$ fluid density, lb/ft³
 - ε = absolute roughness of pipe wall, ft
 - μ = fluid viscosity, cP

COMPRESSIBLE FLUID FLOW IN PIPES

The flow of compressible fluids (e.g., gases and vapors) through pipelines and other restrictions is often affected by changing conditions of pressure, temperature, and physical properties. The densities of gases and vapors vary with temperature and pressure. During isothermal flow, i.e., constant temperature (PV = constant) density varies with pressure. Conversely, in adiabatic flow, i.e., no heat loss (PV^k = constant), a decrease in temperature occurs when pressure decreases, resulting in a density increase. At high pressures and temperatures, the compressibility factor can be less than unity, which results in an increase in the fluid density.

The condition of high ΔP in compressible flow frequently occurs in venting systems, vacuum distillation equipment, and long pipelines [8]. Some design situations involve vapor flows at very high velocities resulting in $\Delta P > 10\%$ of the upstream pressure. Such cases are vapor expanding through a valve, high speed vapor flows in narrow pipes, and vapors flowing in process lines under vacuum conditions. In many cases, ΔP is critical and requires accurate analysis and design. For instance, the inlet pipe ΔP of a safety relief valve should not exceed 3% of the relief valve set pressure (guage) at its relieving capacity for stable operation. This limit is to prevent the rapid opening and closing of the valve, a phenomenom known as chattering, resulting in lowered fluid capacity and subsequent damage of the value seating surfaces. Conversely, the tailpipe or ventline of a relief valve should be designed in such a way that $\Delta P < 10\%$ of the relieving pressure, i.e., set pressure + over pressure in gauge. Figure 3-3 shows a typical tail pipe and relief valve connected to a heat exchanger.

Maximum Flow and Pressure Drop

Determining the maximum fluid flow rate or pressure drop for process design often has the dominant influence on density. As pressure decreases due to piping and component resistance, the gas expands and its velocity increases. A limit is reached when the gas or velocity cannot exceed the sonic or critical velocity. Even if the downstream pressure is lower than the pressure required to reach sonic velocity, the flow rate will still not increase above that evaluated at the critical velocity. Therefore, for a given ΔP , the mass discharge rate through a pipeline is greater for an adiabatic condition (i.e., insulated pipes, where heat transfer is

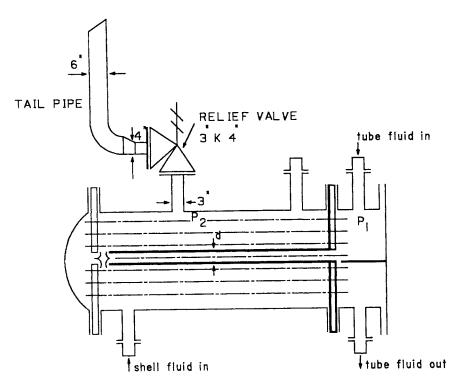


Figure 3-3. Fluid flow through a heat exchanger, relief valve, and tail pipe.

nil) than the rate for an isothermal condition by as much as 20%. There is, however, no difference if the pipeline is more than 1,000 pipe diameters long [10]. In practice, the actual flows are between the two conditions, and the inflow rates are often below 20% even for lines less than 1,000 pipe diameters.

Critical or Sonic Flow and the Mach Number

The flow rate of a compressible fluid in a pipe with a given upstream pressure will approach a certain maximum rate that it cannot exceed even with reduced downstream pressure. The maximum velocity is limited by the velocity of propagation of a pressure wave that travels at the speed of sound in the fluid. The maximum velocity that a compressible fluid can attain in a pipe is known as the sonic velocity, V_s , and can be expressed as:

$$V_{s} = 223 \left(\frac{kT}{M_{w}}\right)^{0.5}, \quad \text{ft/s}$$
$$= 68.1 \left(\frac{kP_{1}}{\rho_{1}}\right)^{0.5}, \quad \text{ft/s}$$
(3-21)

With a high velocity vapor flow, the possibility of attaining critical or sonic flow conditions in a process pipe should be investigated. These occur whenever the resulting pressure drop approaches the following values of ΔP as a percentage of the upstream pressure [11]:

- 1. saturated steam, $\Delta P = 42\%$
- 2. diatomic gases, (e.g., H_2 , N_2 , O_2), $\Delta P = 47\%$
- 3. triatomic and higher molecular weight gases including hydrocarbon vapors and superheated steam, $\Delta P = 45\%$. Vapor flow at or near this maximum velocity should be avoided because a critical pressure, P_c, is attained at the sonic velocity and any ΔP beyond P_c will translate into shock waves and critical turbulence instead of being converted into useful kinetic energy.

In the case of a high pressure header, the flow may be sonic at the exit. Therefore, it is often necessary to check that the outlet pressure of each pipe segment is not critical. If P_c is less than terminal P_2 , the flow is subcritical. If however, P_c is greater than P_2 , then the flow is critical.

Although, it may be impractical to keep the flow in high pressure subheaders below sonic, Mak [12] suggests that the main flare header should not be sized for critical flow at the outlet of the flare stack. This would obviate the undesirable noise and vibration resulting from sonic flow. Crocker's [13] equation for critical pressure can be expressed as:

$$P_{c} = \frac{G}{11400d^{2}} \left(\frac{RT}{k(k+1)}\right)^{0.5}$$
, psia (3-22)

where $R = (1544/29 \cdot SpGr)$, molar gas constant

SpGr = molecular weight of the gas ÷ molecular weight of air

The upstream fluid velocity is:

$$V = \frac{0.0509G}{d^2 \rho_1}, \quad \text{ft/s}$$
 (3-23)

A recommended compressible fluid velocity for trouble-free operation is $V \le 0.6V_s$. The design criteria for compressible fluid process lines (e.g., carbon steel) are shown in Table 3-4.

The Mach Number

The Mach number, M, is the velocity of the gas divided by the velocity of the sound in the gas and can be expressed as:

$$M = \frac{V}{V_s}$$
(3-24)

The exit Mach number for compressible isothermal fluid has been shown to be $M \neq 1$, but $1/\sqrt{k}$, where k is the ratio of the fluid specific heat capacities at constant pressure to that at constant volume. Table 3-5 shows the k values for some common gases. The following cases are such:

- 1. for $M < 1/\sqrt{k}$, the flow is subsonic
- 2. for $M = 1/\sqrt{k}$, the flow is sonic
- 3. for $M > 1/\sqrt{k}$, the flow is supersonic

Case 3 is produced under certain operating conditions in the throttling process (e.g., a reduction in the flow cross-sectional area). Kirkpatrick [14] indicates that there is a maximum length for which continuous flow is applied for an isothermal condition, and this corresponds to

Table 3-4Recommended Fluid Velocity and Maximum△P for Carbon Steel Vapor Lines

Turne of Coursian	Recommended	Maximum ∆P
Type of Service	Velocity ft/sec.	psi/100 ft.
1. General Recommendation		
Fluid pressure psig		
Subatmospheric		0.18
$0 < P \le 50$		0.15
$50 < P \le 150$		0.30
$150 < P \le 200$		0.35
$200 < P \le 500$		1.0
P > 500		2.0
2. Tower Overhead		
Pressure ($P > 50$ psia)	40-50	0.2-0.5
Atmospheric	60-100	
Vacuum (P < 10 psia)		0.05-0.1
3. Compressor Piping Suction	75-200	0.5
4. Compressor Piping Discharge	100-250	1.0
Gas lines with battery limits.		0.5
Refrigerant Suction lines	15-35	
Refrigerant Discharge lines	35-60	
Steam lines		
1. General Recommendation		
Maximum: Saturated	200	
Superheated	250	
Steam pressure in psig.		
0–50	167	0.25
50-150	117	0.40
150-300		1.0
>300		1.5
2. High pressure Steam Lines		
short (L < 600 ft)		1.0
Long (L > 600 ft)		0.5
3. Exhaust Steam lines		0.5
(P > atmosphere)		0.5
Leads to Exhaust Header		1.5
4. Relief valve discharge	0.5V.	
Relief valve, Entry point at silencer	V _s	

Gas	Chemical Formula or Symbol	Approximate Molecular Weight	k(Cp/C _v)
Acetylene (Ethyne)	C_2H_2	26.0	1.30
Air	_	29.0	1.40
Ammonia	NH ₃	17.0	1.32
Argon	A	39.9	1.67
Butane	$C_4 H_{10}$	58.1	1.11
Carbon Dioxide	CO ₂	44.0	1.30
Carobon Monoxide	CO	28.0	1.40
Chlorine	C1 ₂	70.9	1.33
Ethane	$C_2 \tilde{H}_6$	30.0	1.22
Ethylene	C_2H_4	28.0	1.22
Helium	He	4.0	1.66
Hydrogen Chloride	HC1	36.5	1.41
Hydrogen	Η,	2.0	1.41
Methane	\tilde{CH}_4	16.0	1.32
Methyl Chloride	CH ₃ C1	50.5	1.20
Natural Gas		19.5	1.27
Nitric Oxide	NO	30.0	1.40
Nitrogen	N_2	28.0	1.41
Nitrous Oxide	N_2O	44.0	1.31
Oxygen	O_2	32.0	1.40
Propane	$\overline{C_3H_8}$	44.1	1.15
Propylene (Propene)	C_3H_6	42.1	1.14
Sulfur Dioxide	SO_2	64.1	1.26

Table 3-5Approximate k values for some common gases (68°F, 14.7 psia)

 $M = 1/\sqrt{k}$. The limitation for isothermal flow, however, is the heat transfer required to maintain a constant temperature. Therefore, when $M < 1/\sqrt{k}$, heat must be added to the stream to maintain constant temperature. For $M < \sqrt{k}$, heat must be rejected from the stream. Depending on the ratio of specific heats, either condition could occur with subsonic flow. Therefore, to maintain isothermal flow during heat transfer, high temperatures require high Mach numbers and low temperatures require low Mach numbers.

Mathematical Model of Compressible Isothermal Flow

The derivations of the maximum flow rate and pressure drop of compressible isothermal flow are based on the following assumptions:

- 1. isothermal compressible fluid
- 2. no mechanical work done on or by the system
- 3. the gas obeys the perfect gas laws
- 4. a constant friction factor along the pipe
- 5. steady state flow or discharge unchanged with time

Figure 3-4 illustrates the distribution of fluid energy with work done by the pump and heat added to the system. Table 3-6 gives friction factors for clean commercial steel pipes with flow in zones of complete turbulence.

Flow Rate Through Pipeline

Bernoulli's equation for the steady flow of a fluid can be expressed as:

$$\int_{1}^{2} \frac{\mathrm{dP}}{\rho} + \frac{V^{2}}{2g_{c}\alpha} + \frac{g}{g_{c}}\Delta Z + h_{L} + \delta W_{s} = 0 \qquad (3-25)$$

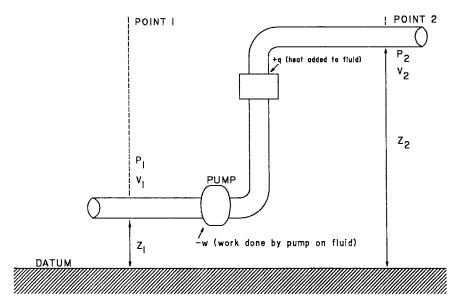


Figure 3-4. Energy aspects of a single-stream piping system.

Nominal Size (inch)	Friction Factor (f)
1	0.023
1.5	0.0205
2	0.0195
3	0.0178
4	0.0165
5	0.016
6	0.0152
8	0.0142
10	0.0136
12	0.0132
14	0.0125
16	0.0122
18	0.12
20	0.0118
24	0.0116

Table 3-6Friction Factors for Total Turbulence in New Commerical SteelPipes

where α (the dimensionless velocity distribution) = 1 for turbulent or plug flow. Assuming no shaft work is done (i.e., $\delta W_s = 0$), then Equation 3-25 becomes

$$\frac{P_2 - P_1}{\rho} + \frac{V_2^2 - V_1^2}{2g_c\alpha} + \frac{g}{g_c}(Z_2 - Z_1) + h_L = 0 \qquad (3-26)$$

For expanding gas flow, $V_2 \neq V_1$; with horizontal pipe, $Z_2 = Z_1$. Hence, the differential form of Bernoulli's equation can be expressed as

$$-\frac{\mathrm{dP}}{\rho} = \frac{1}{g_{\rm c}} \,\mathrm{V}\mathrm{dV} + \mathrm{h_L} \tag{3-27}$$

where ρ is constant and the velocity head is

$$h_{L} = K \frac{V^2}{2g_c}$$
(3-28)

The mass flow rate through the pipe is $G = \rho VA$ or

$$\frac{G}{A} = \rho V = constant$$
 (3-29)

Because both density and velocity change along the pipeline, Equation 3-29 can be expressed in differential form as:

$$\rho dv + V d\rho = 0 \tag{3-30}$$

In expanding gas flow, the pressure and density ratio are constant:

$$\frac{P}{\rho} = \frac{dP}{d\rho}$$
(3-31)

and, with respect to initial condition, P_1 and ρ_1 ,

$$\frac{P}{\rho} = \frac{dP}{d\rho} = \frac{P_1}{\rho_1}$$
(3-32)

From Equation 3-30

$$dV = -\frac{V}{\rho}d\rho \tag{3-33}$$

and

$$\frac{\mathrm{dP}}{\mathrm{P}} = \frac{\mathrm{d\rho}}{\mathrm{\rho}} \tag{3-34}$$

Therefore

$$\frac{1}{\rho} = \frac{1}{P} \left(\frac{dP}{d\rho} \right) = \frac{1}{P} \left(\frac{P_1}{\rho_1} \right)$$
(3-35)

Substituting Equation 3-34 into Equation 3-33

$$dV = -\frac{V}{\rho} \left(\frac{dP}{P}\right) \rho \tag{3-36}$$

Therefore

$$dV = -V\left(\frac{dP}{P}\right)$$
(3-37)

Substituting Equation 3-37 into Equation 3-27

$$-\frac{\mathrm{dP}}{\rho} = \frac{1}{\mathrm{g}_{\mathrm{c}}} \,\mathrm{V}\left(-\mathrm{V}\,\frac{\mathrm{dP}}{\mathrm{P}}\right) + \frac{\mathrm{K}\mathrm{V}^{2}}{2\mathrm{g}_{\mathrm{c}}} \tag{3-38}$$

$$-\frac{\mathrm{dP}}{\rho} = -\frac{1}{g_{\mathrm{c}}} \, \mathrm{V}^2 \left(\frac{\mathrm{dP}}{\mathrm{P}}\right) + \frac{\mathrm{KV}^2}{2g_{\mathrm{c}}} \tag{3-39}$$

Therefore,

$$-dP = \frac{V^2}{g_c} \rho \left(\frac{K}{2} - \frac{dP}{P}\right)$$
(3-40)

From the mass flow rate, $G = \rho VA$

$$V = \frac{G}{\rho A}$$
(3-41)

Substituting Equation 3-41 into Equation 3-40

$$-dP = \frac{G^2}{g_c A^2 \rho^2} \bullet \rho \left(\frac{K}{2} - \frac{dP}{P}\right)$$
(3-42)

In terms of the initial conditions of pressure and density, i.e., substituting Equation 3-35 into Equation 3-42

$$-dP = \left(\frac{G}{A}\right)^{2} \frac{1}{g_{c}} \left\{\frac{1}{P}\left(\frac{P_{1}}{\rho_{1}}\right)\right\} \left[\frac{K}{2} - \frac{dP}{P}\right]$$
(3-43)

Integrating Equation 3-43 gives:

$$-2\int_{1}^{2} P dP = \left(\frac{G}{A}\right)^{2} \left(\frac{P_{1}}{\rho_{1}}\right) \frac{1}{g_{c}} \left\{K - 2\int_{1}^{2} \frac{dP}{P}\right\}$$
(3-44)

which is

$$P_{1}^{2} - P_{2}^{2} = \left(\frac{G}{A}\right)^{2} \frac{P_{1}}{\rho} \left(\frac{1}{g_{c}}\right) \left\{K + 2\ln\frac{P_{1}}{P_{2}}\right\}$$
(3-45)

Therefore, for maximum flow rate through the pipe,

$$G = \left[\left(\frac{A^2 \rho_1 g_c}{K_{\text{TOTAL}} + 2 \ln \frac{P_1}{P_2}} \right) \left(\frac{P_1^2 - P_2^2}{P_1} \right) \right]^{0.5}$$
(3-46)

where K_{TOTAL} is the total velocity head due to friction, fittings, and valves:

$$K_{TOTAL} = f_D \frac{L}{D} + \sum K_f$$
 (pipe fittings + valves)

 ΣK_f (pipe fittings + valves) is the sum of the pressure loss coefficient for all the fittings and valves in the line. Expressing the maximum fluid rate in pounds per hour, Equation 3-46 becomes

$$G = 1335.6d^{2} \left[\left(\frac{\rho_{1}}{K_{\text{TOTAL}} + 2\ln\frac{P_{1}}{P_{2}}} \right) \left(\frac{P_{1}^{2} - P_{2}^{2}}{P_{1}} \right) \right]^{0.5}$$
(3-47)

Pipeline Pressure Drop

If ΔP due to velocity acceleration is relatively small compared with the frictional drop, then $\ln(P_1/P_2)$ may be neglected. Therefore Equation 3-47 becomes

$$G = 1335.6d^{2} \left[\frac{\rho_{1}}{K_{\text{TOTAL}}} \left(\frac{P_{1}^{2} - P_{2}^{2}}{P_{1}} \right) \right]^{0.5}$$
(3-48)

Putting $C = 1335.6d^2$

$$\frac{G^2}{C^2} = \frac{\rho_1}{K_{\text{TOTAL}}} \bullet \frac{P_1^2 - P_1^2}{P_1}$$
(3-49)

i.e.,

$$P_{1}^{2} - P_{2}^{2} = \frac{P_{1}G^{2}K_{\text{TOTAL}}}{\rho_{1}C^{2}}$$
$$P_{2} = \left[P_{1}^{2} - \frac{P_{1}G^{2}K_{\text{TOTAL}}}{\rho_{1}C^{2}}\right]^{0.5}$$
(3-50)

Therefore, the pressure drop

$$\Delta \mathbf{P} \cong \mathbf{P}_1 - \mathbf{P}_2$$

i.e.,

$$\Delta P \cong P_1 - \left[P_1^2 \frac{P_1 G^2 K_{\text{TOTAL}}}{\rho_1 C^2} \right]^{0.5}, \quad \text{psi}$$
(3-51)

Nomenclature

- A = pipe cross-sectional area, ft^2
- d = pipe internal diameter, inch
- D = pipe internal diameter, ft
- f_{c} = Chen friction factor
- $f_{\rm D}$ = Darcy friction factor
- g = acceleration due to gravity (32ft/s²)
- $g_c = \text{conversion factor} \left(32.174(lb_m/lb_f) \bullet (ft/s^2) \right)$
- G =fluid flow rate lb/h
- k = ratio of specific heat capacities (C_p/C_v)
- K = resistance coefficient due to pipe fittings plus valves
- K_f = resistance coefficient due friction pipe fittings + valves
- K_{Total} = total resistance coefficient
 - L_{st} = length of straight pipe, ft
 - M_w = molecular weight of fluid (lb/lb.mol)

M = Mach number

- N_{Re} = Fluid Reynolds number (6.31G/dµ)
 - P_1 = inlet fluid pressure, psia
 - P_2 = outlet fluid pressure, psia
 - $P_c = critical pressure, psia$
 - $R = molar gas constant (1544/29 \times SpGr.)$
- SpGr = molecular weight of gas ÷ molecular weight of air
 - T =fluid temperature, °F
 - V = fluid velocity, ft/s
 - V_s = fluid sonic or critical velocity, ft/s
 - Z = compressibility factor
 - α = correction factor (1 for turbulent or plug flow)
 - ρ = fluid density, (lb_m/ft³)(PM_w/10.73ZRT)
 - μ = fluid viscosity, cP
 - ε = absolute pipe roughness, ft (0.00015 ft for carbon steel)
 - $\Delta = difference$

Subscripts

1 = upstream point in pipe

2 =downstream point in pipe

TWO-PHASE FLOW IN PROCESS PIPING

Two-phase flow often presents design and operational problems not associated with liquid or gas flow. For example, several different flow patterns may exist along the pipeline. Frictional pressure losses are more difficult to estimate, and in the case of a cross-country pipeline, a terrain profile is necessary to predict pressure drops due to elevation changes. The downstream end of a pipeline often requires a separator to separate the liquid and vapor phases, and a slug catcher may be required to remove liquid slugs.

Static pressure losses in gas-liquid flow differ from those in singlephase flow because an interface can be either smooth or rough, depending on the flow pattern. Two-phase pressure losses may be up to a factor of 10 higher than those in single-phase flow. In the former, the two phases tend to separate and the liquid lags behind. Most published correlations for two-phase pressure drop are empirical, and therefore, limited by the range of data for which they were derived [15,16,17]. If two-phase situations are not detected, pressure drop problems may develop that can prevent systems from operating.

Flow Patterns

In determining the type of flow in a process pipeline, designers refer to a diagram similar to Figure 3-5, which is known as the Baker map. Figure 3-6 shows the types of flow regimes that can exist in a horizontal

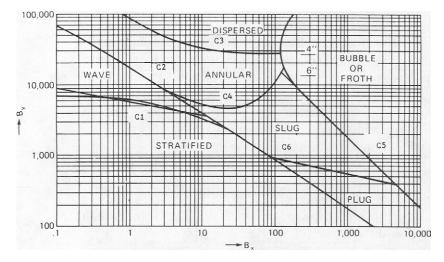


Figure 3-5. Baker parameters for two-phase flow regimes with modified boundaries. Source: Baker [15].

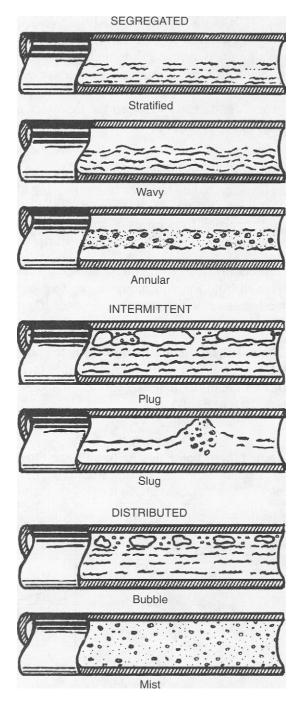


Figure 3-6. Flow regimes in horizontal two-phase flow.

pipe, and Table 3-7 lists the characteristic linear velocities of the gas and liquid phases in each flow regime. Seven types of flow patterns are considered in evaluating two-phase flow, and only one type can exist in a pipeline at a time. But as conditions change (e.g., velocity, roughness, and elevation), the type of flow pattern may also change. The pressure drop can also vary significantly between the flow regimes. The seven types of flow regimes in order of increasing gas rate at a constant liquid flow rate are:

Bubble or Froth Flow. Bubbles of gas are dispersed throughout the liquid and are characterized by bubbles of gas moving along the upper part of the pipe at approximately the same velocity as the liquid. It occurs for liquid superficial velocities of about 5-15 ft/sec (1.5 to 4.5 m/sec) and gas superficial velocities of about 1-10 ft/s (0.3 to 3 m/sec).

Plug Flow. Alternate plugs of liquid and gas move along the upper part of the pipe and liquid moves along the bottom of the pipe. Plug flow occurs for liquid velocities less than 2 ft/sec (0.6m/sec) and gas velocities less than about 3 ft/sec (0.9 m/sec).

Stratified Flow. The liquid phase flows along the bottom of the pipe while the gas flows over a smooth liquid-gas interface. It occurs for liquid velocities less than 0.5 ft/sec (0.15 m/sec) and gas velocities of about 2-10 ft/sec (0.6 to 3 m/s).

Regime	Liquid phase ft/s	Vapor phase ft/s
Bubble or Froth	5–15	0.5–2
Plug	2	<4
Stratified	<0.5	0.5–10
Wave	<1.0	>15
Slug	15 (but less than vapor velocity)	3–50
Annular	<0.5	>20
Dispersed, Spray or Mist	close to vapor velocity	>200

 Table 3-7

 Characteristic Linear Velocities of Two-Phase Flow Regimes

Wave Flow. Wave flow is similar to stratified flow except that the gas is moving at a higher velocity and the gas-liquid interface is distributed by waves moving in the direction of flow. It occurs for liquid velocities less than 1 ft/sec (0.3 m/sec) and gas velocities from about 15 ft/sec. (4.5 m/sec).

Slug Flow. This pattern occurs when waves are picked up periodically by the more rapidly moving gas. These form frothy slugs that move along the pipeline at a much higher velocity than the average liquid velocity. This type of flow causes severe and, in most cases, dangerous vibrations in equipment because of the high velocity slugs against fittings.

Annular Flow. In annular flow, liquid forms around the inside wall of the pipe and gas flows at a high velocity through the central core. It occurs for gas velocities greater than 20 ft/sec (6 m/sec).

Dispersed, Spray, or Mist Flow. Here, all of the liquid is entrained as fine droplets by the gas phase. Dispersed flow occurs for gas velocities greater than 200 ft/sec (60 m/sec).

Flow Regimes

Establishing the flow regime involves determining the Baker parameters B_x and B_y from the two-phase system's characteristics and physical properties. The Baker parameters can be expressed as:

$$\mathbf{B}_{x} = 53 \left[\frac{\mathbf{W}_{L}}{\mathbf{W}_{G}} \right] \left\{ \frac{(\rho_{L} \rho_{G})^{0.5}}{\rho_{L}^{2/3}} \right\} \left(\frac{\mu_{L}^{1/3}}{\sigma_{L}} \right)$$
(3-52)

$$B_{y} = 2.16 \left(\frac{W_{G}}{A}\right) \frac{1}{(\rho_{L}\rho_{G})^{0.5}}$$
(3-53)

 B_x depends on the weight ratio and the physical properties of the liquid and vapor phases. It is independent of pipe size; therefore, it remains constant once calculated from the characteristics of the liquid and vapor, and its position on the Baker map changes only if the liquid-vapor proportion changes. B_y depends on the vapor-phase flow rate, the vapor and liquid densities, and pipe sizes. The practical significance of the pipe size, however, is the effect on the frictional losses. The point B_x , B_y determines the flow regime for the calculated liquid-vapor ratio and the liquid's and vapor's physical properties. With increasing vapor content, B_x , B_y moves up and to the left on the map.

The boundaries of the various flow pattern regions depend on the mass velocity of the gas phase. These boundaries are represented by analytical equations developed by Coker [18]. These equations are used as the basis for determining the prevailing regime for any given flow rates and physical properties of the liquid and vapor. The mathematical models representing the boundaries of the flow regimes are:

$$C_1: \ln B_y = 9.774459 - 0.6548(\ln B_x)$$
 (3-54)

$$C_2$$
: ln $B_y = 8.67694 - 0.1901(ln B_x)$ (3-55)

$$C_3$$
: ln $B_y = 11.3976 - 0.6084(ln B_x) + 0.0779(ln B_x)^2$ (3-56)

$$C_4$$
: ln $B_y = 10.7448 - 1.6265(ln B_x) + 0.2839(ln B_x)^2$ (3-57)

$$C_5: \ln B_v = 14.569802 - 1.0173(\ln B_x)$$
 (3-58)

$$C_6: \ln B_v = 7.8206 - 0.2189(\ln B_x)$$
 (3-59)

Pressure Drop

Various studies have been conducted in predicting the two-phase frictional pressure losses in pipes. The Lockhart-Martinelli correlations [19] shown in Figure 3-7 are employed. The basis of the correlations is that the two-phase pressure drop is equal to the single-phase pressure drop of either phase multiplied by a factor that is derived from the singlephase pressure drops of the two-phases. The total pressure drop is based on the vapor-phase pressure drop. The pressure drop computation is based on the following assumptions:

- 1. The two-phase flow is isothermal and turbulent in both liquid and vapor phases.
- 2. The pressure loss is less than 10% of the absolute upstream pressure.

The two-phase pressure drop can be expressed as:

$$\frac{\Delta PT}{100 ft} = \frac{\Delta PG}{100 ft} \bullet YG \tag{3-60}$$

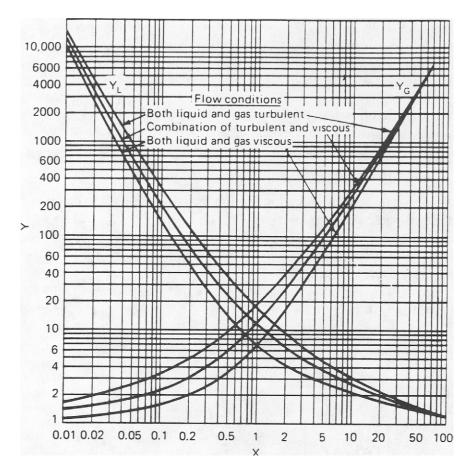


Figure 3-7. Lockhart-Martinelli pressure drop correlation. Source: Lockhart and Martinelli [19].

where YG is the two-phase flow modulus. YG is a function of the Lockhart-Martinelli two-phase flow modulus X, which is defined as:

$$X = \left(\frac{\Delta PL}{\Delta PG}\right)^{0.5}$$
(3-61)
YG = f(x)

The value of YG for the different flow regimes is determined as follows: For bubble or froth flow

$$YG = \left[\frac{14.2X^{0.75}}{(WL/A)^{0.1}}\right]^2$$
(3-62)

For plug flow

$$YG = \left[\frac{27.315X^{0.855}}{(WL/A)^{0.17}}\right]^2$$
(3-63)

For stratified flow

$$YG = \left[\frac{15,400X}{(WL/A)^{0.8}}\right]^2$$
(3-64)

For wave flow, the Huntington friction factor [20] is used to determine the two-phase pressure loss:

$$H_{x} = \left(\frac{HL}{WG}\right) \left(\frac{\mu_{L}}{\mu_{G}}\right)$$
(3-65)

 $\ln FH = 0.2111 \ln(H_x) - 3.993$ (3-66)

$$\frac{\Delta PT}{100 ft} = \frac{0.000336(FH)(WG)^2}{d^5 \rho_G}$$
(3-67)

For slug

$$YG = \left[\frac{1190X^{0.815}}{(WL/A)^{0.5}}\right]^2$$
(3-68)

For annular flow

 $YG = (aX^{b})^{2}$ (3-69)

$$a = 4.8 - 0.3125d$$

$$b = 0.343 - 0.021d$$

d = pipe inside diameter, inch

$$d = 10$$
 for 12 inch and larger sizes

For dispersed or spray flow

 $C_0 = 1.4659$ $C_1 = 0.49138$ $C_2 = 0.04887$ $C_3 = -0.000349$

Once the two-phase flow modulus (YG) for the particular flow regime has been calculated, Equation 3-60 can be used to calculate Δ PT/100ft, the two-phase, straight-pipe pressure drop. The pressure drop of liquid or gas flowing alone in a straight pipe can be expressed as:

$$\frac{\Delta PT}{100 ft} = \frac{0.000336 f_D W_x^2}{d^5 \rho_G}, \quad psi/100 ft$$
(3-71)

Equation 3-71 can be modified for calculating the overall pressure drop of the two-phase flow for the total length of pipe plus fittings, L from Equation 3-5, based on the gas-phase pressure drop.

$$\Delta PT_{overall} = \frac{0.000336 \bullet f_{D} \bullet WG^{2} \bullet YG \bullet L}{100d^{5}\rho_{G}}, \quad psi \qquad (3-72)$$

The velocity of the two-phase fluid is

$$V = \frac{0.0509}{d^2} \left[\frac{WG}{\rho_G} + \frac{WL}{\rho_L} \right]$$
(3-73)

Erosion-Corrosion

Depending on the flow regime, the liquid in a two-phase flow system can be accelerated to velocities approaching or exceeding the vapor velocity. In some cases, these velocities are higher than what would be desirable for process piping. Such high velocities lead to a phenomenon known as erosion-corrosion, where the corrosion rate of a material is accelerated by an erosive material or force (in this case, the high-velocity liquid).

An index [21] based on velocity head can indicate whether erosioncorrosion may become significant at a particular velocity. It can be used to determine the range of mixture densities and velocities below which erosion-corrosion should not occur. This index is:

$$\rho_{\rm M} U_{\rm M}^2 \le 10,000 \tag{3-74}$$

where the mixture density is:

$$\rho_{\rm M} = \frac{WL + WG}{\left(\frac{WL}{\rho_{\rm L}} + \frac{WG}{\rho_{\rm G}}\right)}$$
(3-75)

and the mixture velocity is:

$$U_{M} = U_{G} + U_{L}$$
$$= \left(\frac{WG}{3600\rho_{G}A} + \frac{WL}{3600\rho_{L}A}\right)$$
(3-76)

Nomenclature

A = inside cross-sectional area of pipe, ft^2

- B_x , B_y = Baker parameters for determining for regime
 - d = pipe inside diameter, inch
 - D = pipe inside diameter, ft
 - f_{c} = Chen friction factor
 - $f_D = Darcy friction factor$
 - $f_F = Fanning friction factor$
 - FH = Huntington friction factor
 - H_x = Huntington flow factor
 - ID = internal diameter of pipe, inch
 - K = excess head loss for a fitting, velocity heads
 - $K_1 = K$ for fitting at $N_{Re} = 1$ velocity heads
 - $K\infty = K$ for large fittings at $N_{Re} = \infty$ velocity heads
 - L = total fitting equivalent pipe length plus straight pipe length, ft
 - L_{eq} = equivalent length of pipe due to fittings, ft
 - L_{st} = straight pipe length, ft
 - L_{Total} = total length of pipe, ft
 - MW = molecular weight
 - N_{Re} = Reynolds number
 - $N_{ReL} =$ liquid Reynolds number
 - $N_{ReG} = gas Reynolds number$

 $\Delta PL/100$ ft = pressure drop of liquid if flowing alone in the pipe (psi/100 ft)

 $\Delta PG/100$ ft =pressure drop of gas if flowing alone in the pipe (psi/100 ft)

- $\Delta PT/100ft$ = pressure drop of the two-phase mixture in straight pipe (psi/100 ft)
 - $\Delta PT_{overall}$ = overall pressure drop of the two-phase mixture for the total length plus fittings, psi

 $U_{L} =$ liquid velocity, ft/s

 $U_{G} = gas velocity, ft/s$

 U_{M} = velocity of the two-phase mixture, ft/s

V = velocity of fluid in a pipe, ft/s

WL = liquid flow rate, lb/h

WG = gas flow rate, lb/h

- $W_x =$ flow rate of either liquid or gas, lb/h
 - X = Lockhart-Martinelli two-phase flow modulus $(\Delta PL/\Delta PG)^{0.5}$
- Y_G = two-phase flow modulus
 - ε = absolute roughness of pipe wall, ft (0.00015 ft for carbon steel)
- $\rho_{\rm L}$ = liquid density, lb/ft³

 $\rho_{\rm G}$ = gas density, lb/ft³

- $\rho_{\rm M}$ = density of liquid-gas mixture, lb/ft³
- μ_x = viscosity of liquid or gas, cP
- $\mu_{\rm L}$ = liquid viscosity, cP
- μ_{G} = gas viscosity, cP
- $\sigma_{\rm L}$ = surface tension of liquid (dyne/cm)

VAPOR-LIQUID TWO-PHASE VERTICAL DOWNFLOW

An understanding of two-phase flow is necessary for sound piping design because almost all chemical process plants encounter two-phase flow conditions. Two-phase vertical downflow presents its own problems in horizontal flow. In a vertical flow, large vapor bubbles, known as slug flow, are formed in the liquid stream. This flow regime is associated with pipe vibration and pressure pulsation. With bubbles greater than 1 inch in diameter and the liquid viscosity less than 100 cP, slug flow region can be represented by dimensionless numbers for liquid and vapor phases respectively (Froude numbers (N_{Fr})_L and (N_{Fr})_G). These are related by the ratio of inertial to gravitational force and are expressed as:

$$(N_{Fr})_{L} = \frac{VL}{(gD)^{0.5}} \left(\frac{\rho_{L}}{\rho_{L} - \rho_{G}}\right)^{0.5}$$
(3-77)

$$(N_{Fr})_{G} = \frac{VG}{(gD)^{0.5}} \left(\frac{\rho_{L}}{\rho_{L} - \rho_{G}}\right)^{0.5}$$
(3-78)

The velocities VG and VL are superficial velocities based on the total pipe cross section. These Froude numbers exhibit several features in the range $0 < (N_{Fr})_L$ and $(N_{Fr})_G < 2$.

Simpson [3] illustrates the values of $(N_{Fr})_L$ and $(N_{Fr})_G$ with water flowing at an increased rate from the top of an empty vertical pipe. As the flow rate further increases to the value $(N_{Fr})_L = 2$, the pipe floods and the total cross section is filled with water. If the pipe outlet is further submerged in water and the procedure is repeated, long bubbles will be trapped in the pipe below $(N_{Fr})_L = 0.31$. However, above $(N_{Fr})_L = 0.31$, the bubbles will be swept downward and out of the pipe.

If large long bubbles are trapped in a pipe (d ≥ 1 inch) in vertically down flowing liquid having a viscosity less than 100 cP and the Froude number for liquid phase, $(N_{Fr})_L \leq 0.3$, the bubbles will rise. At higher Froude numbers, the bubbles will be swept downward and out of the pipe. A continuous supply of vapor causes the Froude number in the range $0.31 \leq (N_{Fr})_L < 1$ to produce pressure pulsations and vibration. These anomalies are detrimental to the pipe and must be avoided. If the Froude number is greater than 1.0, the frictional force offsets the effect of gravity, and thus requires no pressure gradient in the vertical downflow liquid. This latter condition depends on the Reynolds number and pipe roughness.

The Equations

The following equations will calculate Froude numbers for both the liquid and gas phases. The computer program will print out a message indicating whether the vertical pipe is self-venting, whether pulsating flow occurs, or whether no pressure gradient is required.

$$D = \frac{d}{12}, \quad ft \tag{3-79}$$

Area =
$$\frac{\pi D^2}{4}$$
, ft (3-80)

$$VL = \frac{WL}{3600 \bullet \rho_L \bullet Area}, \quad ft/s$$
 (3-81)

$$VG = \frac{WG}{3600 \bullet \rho_G \bullet Area}, \quad ft/s$$
 (3-82)

$$FRNL = \frac{VL}{(gD)^{0.5}} \left[\frac{\rho_L}{\rho_L - \rho_G} \right]^{0.5}$$
(3-83)

FRNG =
$$\frac{VG}{(gD)^{0.5}} \left[\frac{\rho_G}{\rho_L - \rho_G} \right]^{0.5}$$
 (3-84)

The Algorithm

If FRNL < 0.31, vertical pipe is SELF-VENTING

else

 $0.3 \le FRNL < 1.0$, PULSE FLOW, and may result in pipe vibration FRNL > 1.0, NO PRESSURE GRADIENT

Nomenclature

 $\begin{array}{l} \mbox{Area}=\mbox{inside cross-sectional area of pipe, ft}^2\\ \mbox{d}=\mbox{inside diameter of pipe, inch}\\ \mbox{D}=\mbox{inside diameter of pipe, ft}\\ \mbox{FRNL, } (N_{Fr})_L = \mbox{Froude number of liquid phase, dimensionless}\\ \mbox{FRNG, } (N_{Fr})_G = \mbox{Froude number of vapor phase, dimensionless}\\ \mbox{g}=\mbox{gravitational constant, } 32.2 \mbox{ ft/s}^2\\ \mbox{VL} = \mbox{liquid velocity, ft/s}\\ \mbox{VG} = \mbox{vapor velocity, ft/s}\\ \mbox{WL} = \mbox{liquid flowrate, lb/h}\\ \mbox{WG} = \mbox{vapor flowrate, lb/h}\\ \mbox{\rho}_L = \mbox{liquid density, lb/ft}^3\\ \mbox{\rho}_G = \mbox{vapor densit, lb/ft}^3 \end{array}$

LINE SIZES FOR FLASHING STEAM CONDENSATE

When a liquid is flowing near its saturation point (i.e., the equilibrium or boiling point) in a pipe line, decreased pressure will cause vaporization. The higher the pressure difference, the greater the vaporization resulting in flashing of the liquid. Steam condensate lines cause a two-phase flow condition, with hot condensate flowing to a lower pressure through short and long lines. For small lengths with low pressure drops, and the outlet end being a few pounds per square inch of the inlet, the flash will be assumed as a small percentage. Consequently, the line can be sized as an all liquid line. However, caution must be exercised because 5% flashing can develop an important impact on the pressure drop of the system [22].

Sizing of flashing steam condensate return lines requires techniques that calculate pressure drop of two-phase flow correlations. Many correlations have been presented in the literature [15,16,19,23]. Most flow patterns for steam condensate headers fall within the annular or dispersed region on the Baker map. Sometimes, they can fall within the slug flow region; however, the flashed steam in steam condensate lines is less than 30% by weight.

Ruskin [24] developed a method for calculating pressure drop of flashing condensate. Ruskin's method gave pressure drops comparable to those computed by the two-phase flow with good agreement with experimental data. The method employed here is based on a similar technique given by Ruskin. The pressure drop for flashing steam uses the average density of the resulting liquid-vapor mixture after flashing. In addition, the friction factor used is valid for complete turbulent flows in both commercial steel and wrought iron pipe. The pressure drop assumes that the vapor-liquid mixture throughout the condensate line is represented by mixture conditions near the end of the line. This assumption is valid because most condensate lines are sized for low pressure drop, with flashing occurring at the steam trap or valve close to the pipe entrance. If the condensate line is sized for a higher pressure drop, an iterative method must be used. For this case, the computations start at the end of the pipeline and proceed to the steam trap.

The method employed determines the following:

- 1. the amount of condensate flashed for any given condensate header from 15 to 140 psia. Initial steam pressure may vary from 40 to 165 psia.
- 2. the return condensate header temperature.
- 3. the pressure drop (psi/100 ft) of the steam condensate mixture in the return header.
- 4. the velocity of the steam condensate mixture and gives a warning message if the velocity is greater than 5000 ft/min, because this may present problems to the piping system.

The Equations

The following equations are used to determine the pressure drop for flashed condensate mixture [20].

$$WFRFL = B(\ln P_c)^2 - A$$
(3-85)

where

$$A = 0.00671(\ln P_{\rm h})^{2.27}$$
(3-86)

$$\mathbf{B} = \mathbf{e}^{\mathbf{x}} \bullet 10^{-4} + 0.0088 \tag{3-87}$$

and

$$X = 6.122 - \left(\frac{16.919}{\ln P_{h}}\right)$$
(3-88)

$$WG = WFRFL \bullet W \tag{3-89}$$

$$WL = W - WG \tag{3-90}$$

$$TFL = 115.68(P_h)^{0.226}$$
(3-91)

$$\rho_{\rm G} = 0.0029 P_{\rm h}^{0.938} \tag{3-92}$$

$$\rho_{\rm L} = 60.827 - 0.078 P_{\rm h} + 0.00048 P_{\rm h}^2 - 0.0000013 P_{\rm h}^3 \qquad (3-93)$$

$$\rho_{\rm M} = \frac{\rm WG + \rm WL}{\left(\frac{\rm WG}{\rho_{\rm G}} + \frac{\rm WL}{\rho_{\rm L}}\right)}$$
(3-94)

For turbulent flow

$$f = \frac{0.25}{\left[-\log\left(\frac{0.000486}{d}\right)\right]^2}$$
(3-95)

where d = pipe diameter, inch

Pressure drop

$$\Delta PT = \frac{0.000336f \bullet W^2}{d^5 \bullet \rho_M}$$
(3-96)

$$V = \frac{3.054}{d^2} \left[\frac{WG}{\rho_G} + \frac{WL}{\rho_L} \right]$$
(3-97)

If $V \ge 5000$ ft/min, print a warning message as condensate may cause deterioration of the process pipeline.

Nomenclature

d = internal pipe diameter, inch

f = friction factor, dimensionless

- P_c = steam condensate pressure before flashing, psia
- P_{h} = flashed condensate header pressure, psia
- $\dot{\mathbf{V}}$ = velocity of flashed condensate mixture, ft/min
- W = total flow of mixture in condensate header, lb/h

WG = flashed steam flow rate, lb/h

WL = flashed condensate liquid flow rate, lb/h

WFRFL = weight fraction of condensate flashed to vapor

TFL = temperature of flashed condensate, °F

 ΔPT = pressure drop of flashed condensate mixture, psi/100 ft

 ρ_G = flashed steam density, lb/ft³

 $\rho_{\rm I}$ = flashed condensate liquid density, lb/ft³

 $\rho_{\rm M}$ = density of mixture (flashed condensate/steam), lb/ft³

FLOW THROUGH PACKED BEDS

Flow of fluids through packed beds of granular particles occurs frequently in chemical processes. Examples are flow through a fixed-bed catalytic reactor, flow through a filter cake, and flow through an absorption or adsorption column. An understanding of flow through packed beds is also important in the study of sedimentation and fluidization.

An essential factor that influences the design and operation of a dynamic catalytic or adsorption system is the energy loss (pressure drop). Factors determining the energy loss are many and investigators have made simplifying assumptions or analogies so that they could use some of the general equations. These equations represent the forces exerted by the fluids in motion (molecular, viscous, kinetic, static, etc.) to arrive at a useful expression correlating these factor.

Ergun [25] developed a useful pressure drop equation caused by simultaneous kinetic and viscous energy losses and applicable to all types of flow. Ergun's equation relates the pressure drop per unit of bed depth to dryer or reactor system characteristics, such as, velocity, fluid gravity, viscosity, particle size, shape, surface of the granular solids and void fraction. The original Ergun equation is:

$$\frac{\Delta P}{L}g_{c} = \frac{150(1-\varepsilon)^{2}}{\varepsilon^{3}} \bullet \frac{\mu V}{D_{p}^{2}} + \frac{1.75(1-\varepsilon)}{\varepsilon^{3}} \bullet \frac{GV}{D_{p}}$$
(3-98)

where ΔP = pressure drop

- L = unit depth of packed bed
- g_c = dimensional constant 32.174(lb_m/lb_f)(ft/s²)
- μ = viscosity of fluid, lb/ft.h
- V = superficial fluid velocity, ft/s
- D_p = effective particle diameter, ft
 - $\dot{\varepsilon}$ = void fraction of bed
 - ρ = fluid density, lb/ft³

Equation 3-98 gives the total energy loss in fixed beds as the sum of viscous energy loss (the first term on the right side of the equation) and the kinetic or turbulent energy loss (the second term on the right side of the equation). For gas systems, approximately 80% of the energy loss is dependent on turbulence and can be represented by the second term of Equation 3-98. In liquid systems, the viscous term is the major factor.

The Equation

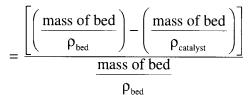
The original Ergun equation of the total energy loss can be rearranged as follows:

$$\Delta PT = \frac{BL}{144} \bullet \frac{1-\varepsilon}{\varepsilon^3} \bullet \frac{G^2}{D_p g_c \rho} \left[\frac{150(2.419\mu)(1-\varepsilon)}{D_p G} + 1.75 \right] (3-99)$$

 ε = fraction void volume of bed, ft³,

void/ft³ of packed tower volume

 $= \frac{\text{volume of void}}{\text{volume of bed}}$



$$=\frac{\rho_c - \rho_b}{\rho_c} \tag{3-100}$$

$$\rho_c = \text{density of catalyst, } \text{lb/ft}^3$$

$$\rho_{\rm b}$$
 = density of the packed bed, lb/ft³

$$\rho = \frac{M_{w}P}{10.73ZT}$$
(3-101)

$$D_{p} = \frac{6(1-\varepsilon)}{S}$$
(3-102)

$$S = \frac{(1-\varepsilon)}{V_{p}} \bullet A_{p}$$
(3-103)

For cylindrical particles

$$V_{p} = \frac{1}{1728} \left[PD^{2} \bullet \frac{\pi}{4} \bullet PL \right]$$
(3-104)

$$A_{p} = \frac{1}{144} \left[\frac{\pi \bullet PD^{2}}{2} + \pi \bullet PD \bullet PL \right]$$
(3-105)

Reynolds number

$$N_{Re} = \frac{GD_{p}}{2.419\mu(1-\epsilon)}$$
(3-106)

Friction factor For laminar flow with $N_{Re} < 1$

$$f_{p} = \frac{150}{N_{Re}}$$
(3-107)

For intermediate Reynolds number

$$f_{p} = \frac{150}{N_{Re}} + 1.75$$
(3-108)

For completely turbulent flow, it is assumed that f_p approaches a constant value for all packed beds with the same relative roughness. The constant is found by experiment to be 1.75 [26].

Nomenclature

A = cross-sectional area of bed, ft^2

 $A_p = area of particles, ft^2$

BL = bed length, ft

 $D_n = effective particle diameter, ft$

 $f_n = friction factor$

 $g_{c}^{r} = \text{gravitational constant} = 4.17 \times 10^{8} ((lb_{m}/lb_{f}) \bullet (ft/h^{2}))$

 $G = superficial mass velocity = W/A, lb/h.ft^2$

- MW = molecular weight of fluid
 - N_{Re} = Reynolds number, dimensionless
 - P = fluid pressure, psia
 - PD = particle diameter, inch

PL = particle length, inch

- S = packed bed surface area, ft^2/ft^3 bed
- T =fluid temperature, °R
- V_{n} = volume of particles, ft³
- \dot{W} = fluid flow rate, lb/h
- Z = compressibility factor
- ΔPT = total pressure drop in packed bed, lb/in²
 - ε = fraction voids in packed bed
 - ρ = density of fluid at flowing conditions, lb/ft³
 - μ = fluid viscosity, cP (1cP = 2.419lb/ft h)

SCREEN HANDLING

The code in this book makes use of a suite of subroutines for screen handling that have been developed from those given by Ward and Bromhead [27,28].

In principle, a number of subroutines are invoked by my code, and these in turn invoke other, lower level subroutines. The top level subroutines are called:

GETTXT GETINT GETFLO

and

MENU

GETTXT is perhaps the simplest of these. It uses **ANSI** control sequences to locate the cursor on the screen, to highlight a box for text entry, and through assembly-language-coded calls to an **MS DOS** function, intercepts individual keypresses and builds them into a text string in the text entry box.

The low level functions that use the ANSI codes are coded into

GOTOXY	Cursor location	(FORTRAN)
REVID	Turn reversed video effect on	(ASM)
NORMAL	Restore normal video (no attributes)	(ASM)
SHWTXT	Display string	(FORTRAN)

From these further calls to assembly language routines that call **DOS** functions are made. These routines are:

GETCHR	Get character from keyboard buffer
SHWBYT	Display character string

Analogous routines to **GETTXT** are provided for inputting INTE-GER and FLOATING POINT (REAL) variables, and they are called **GETINT** and **GETFLO**. In principle, they are identical to **GETTXT**, but they perform the additional tasks of intercepting invalid keypresses (e.g., no alphabetic characters are permitted nor can the decimal point be duplicated, etc.) and converting from a text string into a valid number format. A routine **BEEP** is provided to send ASCII 07 (BEL) to the screen, which drives the PC's internal speaker.

The subroutine **MENU** displays a list of mnemonics on the screen at predefined locations. In my code, these mnemonics are usually in a vertical stack, and they are all accompanied by a line of explanatory text per mnemonic. Use of the cursor keys causes a highlighted bar to step through the mnemonics. Pressing the RETURN key selects the option equivalent to the currently selected mnemonic.

In addition to the GETxxx subroutines, I sometimes need to display text or numbers at predefined locations on the screen. The GOTOXY and SHWTXT calls are put together in DISTXT, DISINT, and DISFLO (DIS implying DISplay). Some other routines which are used to create special effects are

BORDER COLORS

BORDER permits me to draw rectangular boxes on the screen using the extended character set line-drawing characters (both single and double). **COLORS** uses the **ANSI** codes to set foreground and background colors. The advantage of using the ANSI screen control system is that it makes many of the jobs involved in screen control simple for the **FORTRAN** programmer, but at a cost: the computer on which the software is run must have **ANSI.SYS** loaded. Under DOS 5 or DOS 6, **ANSI.SYS** can be loaded "high," which prevents it from taking up valuable conventional **DOS** memory. There are various "enhanced **ANSI.SYS**" utilities which are public domain or shareware, including FANSI and NANSI. Another useful utility is ANSI.COM (PC Magazine, Ziff Davies Publishing), which has the advantage of being unloadable if the user wishes to reclaim the small amount of memory that it consumes.

Title Screen and Keyboard Utilities

,		
CSSANDK	SEGMENT	PUBLIC PARA 'CODE' ASSUME CS:CSSANDK
•		
PUBLIC	CLS	
CLS	PROC	FAR
OLO	PUSH	DS
	MOV	AH, 9
	MOV	DH, SEG CLRSTRG
	MOV	DS, DX
	MOV	DX, OFFSET CLSRTRG
	INT	21H
	POP	DS
CLRSTRG	DB	1BH,'[2J\$'
		1011, [200
CLS	ENDP	
	PUBLIC	REVID
REVID	PROC	FAR
	PUSH	DS
	MOV	AH, 9
	MOV	DX, SEG REVSTR
	MOV	DS, DX
	MOV	•
		DX, OFFSET REVSTR
	INT	21H

Fortran Programs for Chemical Process Design

POP	DS
RET DB ENDP	1BH,'[7m\$'
PUBLIC PROC PUSH MOV MOV MOV MOV INT POP RET	NORMAL FAR DS AH, 9 DX, SEG NORSTR DS, DX DX, OFFSET NORSTR 21H DS
DB ENDP	1BH,'[0m\$'
PUBLIC PROC PUSH MOV MOV LDS INT POP RET ENDP	SHWBYT FAR BP AH, 09H BP, SP DX, [BP+6] 21H BP 04H
PUBLIC PROC PUSH MOV INT MOV LES XOR MOV POP RET ENDP	GETCHR FAR BP AH, 08H 21H BP, SP BX, DWORD PTR[BP+6] AH, AH; this zeroes AH ES:[BX], AX; better than MOV BP 04H
	RET DB ENDP PUBLIC PROC PUSH MOV MOV MOV MOV MOV MOV INT POP RET DB ENDP PUBLIC PROC PUSH MOV MOV LDS INT POP RET ENDP PUBLIC PROC PUSH MOV MOV LDS INT POP RET ENDP PUBLIC PROC PUSH MOV MOV LDS INT POP RET ENDP

KBCLEAR	PUBLIC KBO PROC PUSH MOV	CLEAR FAR BP BP, SP
WHILE:	MOV INT OR JZ MOV INT JMP	AH, 0BH 21H AL, AL SHORT RETURN AH, 08H 21H SHORT WHILE
KBCLEAR	POP RET ENDP	BP
CURSOF		CURSOF FAR BP CX, 2000H AH, 1 10H BP
CURSBLK	PUBLIC PROC PUSH MOV MOV INT POP RET ENDP	CURSBLK FAR BP CX, 0007H AH, 1 10H BP
CURSON	PUBLIC PROC PUSH MOV	CURSON FAR BP CX, 0607H

194 Fortran Programs for Chemical Process Design

	MOV INT POP RET	AH, 1 10H BP
CURSON	ENDP	
, CSSANDK	ENDS	
3	END	
		Fortran Subroutines

100	SUBROUTINE CLS WRITE(*,100) CHAR(27)//'[2J' FORMAT(A4/) RETURN END
100	SUBROUTINE REVID WRITE(*,100) CHAR(27)//'[7m' FORMAT(A4/) RETURN END
100	SUBROUTINE NORMAL WRITE(*,100) CHAR(27)//'[0m' FORMAT(A4/) RETURN END
100	SUBROUTINE SHWBYT (STR, NSTR) CHARACTER*(*) STR WRITE(*,100) STR(1:NSTR) FORMAT(A/) RETURN END

PROBLEMS AND SOLUTIONS

Problem 3-1

A piping isometric (Figure 3-8) shows a 6-inch (Schedule 40) line with six 90°LR elbows and two flow-through tees. The actual length of

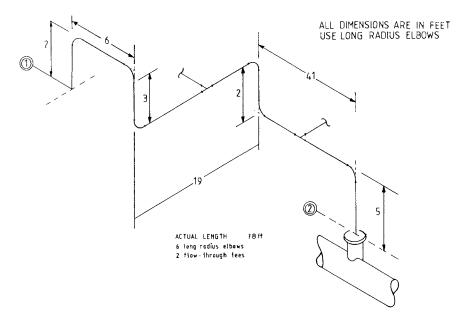


Figure 3-8. Piping isometric layout.

pipe is 78 ft. The fluid flow is 75,000 lb/h and its physical properties are viscosity $\mu = 1.25$ cP, density $\rho = 64.30$ lb/ft³, and pipe roughness $\varepsilon = 0.00015$ ft. Calculate the equivalent length of pipe fittings, total length of the pipe, velocity head, head loss, and the overall pressure drop.

Solution

The computer program PROG31 is a subroutine in the screen handling suites of programs used to calculate the parameters for the 6-inch (Schedule 40) pipe. The software program **PIPECAL** is run to determine the total length of the pipe (i.e., the sum of the straight length of the pipe and the equivalent length due to pipe fittings), the head loss, the pressure drop per 100 ft of pipe, and the overall pressure drop. Table 3-8 gives the computer output from the screen handling software program. The equivalent length due to pipe fittings is 38.3 ft, the total pipe length is 116.3 ft, the excess head loss is 0.196 ft, the pressure drop per 100 ft of the pipe is 0.075 psi, and the overall pressure drop is 0.088 psi.

Table 3-8			
Computer Output From the Screen Handling Programs			

PRESSURE DROP CALCULATION OF AN INCOMPRESSIBLE	
***************************************	**********
PIPE INTERNAL DIAMETER, inch:	6.065
MASS FLOW RATE, 1b/h:	75000.00
VOLUMETRIC FLOW RATE, gal/min.:	145.44
FLUID VISCOSITY, cP:	1.2500
FLUID DENSITY, 1b/ft.^3:	64.3000
FLUID VELOCITY, ft/sec.:	1.614
VELOCITY HEAD LOSS DUE TO FITTINGS:	1.5960
EQUIVALENT LENGTH OF PIPE, ft:	38.3099
ACTUAL LENGTH OF PIPE, ft:	78.0000
TOTAL LENGTH OF PIPE, ft.:	116.3099
REYNOLDS NUMBER:	62424.
PIPE ROUGNESS, ft.:	.000150
DARCY FRICTION FACTOR:	.0211
EXCESS HEAD LOSS, ft:	.1961
PIPE PRESSURE DROP/100 ft, psi/100ft:	.0754
OVERALL PRESSURE DROP OF PIPE, psi:	.0877

Running Pipecal

The computer program **PIPECAL**, as well as the other programs described in the screen handling, has been compiled to calculate the equivalent length due to pipe fittings, the total length of the pipe, velocity head, the head loss, the pressure drop per 100 ft of pipe, and the overall pressure drop for any given pipe size, fluid flow rate, and its physical properties. To run the program from either the floppy diskette or a hard drive, type after the A or C:\> prompt from the command line, PIPECAL.EXE (or PIPECAL) followed by a return key.

For example, A or C:>PIPECAL.EXE (or PIPECAL)

A title is displayed and the user then presses any key from the PC console to continue. Once the screen is cleared, a list of eight options is displayed as follows:

- 1. **OPEN** Open a pre-existing file
- 2. **CREATE** Create a new data file
- 3. **SAVE** Save the data file you just edited
- 4. CALCS Start calculating
- 5. **PRT RES** Print the result file
- 6. **SAVE RES** Save the result file
- 7. **FINISH** Next problem?
- 8 **QUIT** Exit to DOS

PIPECAL is user-friendly and the procedure in running the program is as follows:

- 1. Create a data file (option 2). Assign the value zero to any fitting type that is not included in the calculation. Press a return key or a downward arrow key to continue.
- 2. Calculate the input data (option 4).
- 3. Print the results onto a printer (option 5). The results are also viewed on the screen.
- 4. Quit the program by typing N (option 8) to return to DOS (Disk Operating System).
- 5. Press FINISH (Option 7) before opening or creating a datafile.

Options 1, 3, 6, and 7 can be further used after the datafile has been created.

Before using the software (**PIPECAL**), you must have **ANSI.SYS** installed as a device driver in the **CONFIG.SYS** file. If for example, **ANSI.SYS** is located in the directory **C:\DOS**, the following line must be a part of **CONFIG.SYS**

DEVICE=C:\DOS\ANSI.SYS

Problem 3-2

Calculate the overall pressure drop between points 1 and 2 for a 6-inch (Schedule 40, ID=6.065 inch) line for kerosene. Liquid conditions are: flow rate $Q_{60} = 900$ gpm, density at 60°F, $\rho = 51$ lb/ft³, and temperature, t = 321°F. Figure 3-8 shows an isometric layout of the pipeline with fittings.

Specific gravity at 60°F, $S_{60} = 51/62.37 = 0.82$ Specific gravity at t = 321°F, $S_{321} = 0.72$ Density at t = 321°F, $\rho = 62.37$ (0.72) = 44.9 lb/ft³ Expansion factor, $E = S_{60}/S_{321} = 0.82/0.72 = 1.14$ Flow rate at t = 321°F, $Q = Q_{60}$. E = 900(1.14) = 1026 gpm Viscosity, $\mu = 0.3$ cP Actual length of pipe = 78 ft.

Solution

The computer program PROG32 evaluates the pressure drop for any incompressible fluid per 100 ft of pipe and the overall pressure drop between two points for any given pipe size. The program PROG32 calculates the equivalent length of pipe due to fittings. This is added to the straight length of pipe to give the total length. Table 3-9 gives the input data and computer output for the 6-inch pipe size. The pressure drop per 100 ft of pipe is 1.93 psi, and the overall pressure drop is 3.4 psi.

Table 3-9 Input Data and Computer Output for an Incompressible Fluid in a Pipe Line

G.065 78.0 1026.0 0.0 44.9 0.3 0.0 3.0 0.00015 0.00015 0.00015 PRESSURE DROP CALCULATION OF AN INCOMPRESSIBLE FLUID IN A PIPE LINE ***********************************	DATA32.	DAT					
************************************	44.9	0.3					
MASS FLOW RATE, lb/h.: 369263.30 VOLUMETRIC FLOW RATE, gal/min.: 1026.00 FLUID VISCOSITY, cP: .3000 FLUID DENSITY, lb/ft^3: 44.900 FLUID VELOCITY, ft/sec.: 11.380 VELOCITY HEAD LOSS DUE TO FITTINGS: 3.000 EQUIVALENT LENGTH OF PIPE, ft.: 97.919 ACTUAL LENGTH OF PIPE, ft.: 78.000 TOTAL LENGTH OF PIPE, ft.: 175.919 REYNOLDS NUMBER: 1281127. PIPE ROUGHNESS, ft.: .000150 DARCY FRICTION FACTOR: .0155 EXCESS HEAD LOSS, ft.: 10.867 PIPE PRESSURE DROP /100ft., psi/100ft: 1.9264	PR	ESSURE	DROP CA				**
OVERALL PRESSURE DROP OF PIPE, psi: 3.3889	MAS VOL FLU FLU VEL EQU ACT TOT REY PIP DAR EXC	S FLOW UMETRIC ID VISC ID DENS ID VELC OCITY H IVALEN UAL LENC NOLDS H E ROUGH CY FRIC ESS HEZ	RATE, 1 C FLOW R COSITY, 5 SITY, 1b DCITY, f HEAD LOS T LENGTH OF STH OF P NUMBER: DTION FA AD LOSS,	b/h.: ATE, gal/min.: cP: /ft^3: t/sec.: S DUE TO FITTINGS: OF PIPE, ft.: PIPE, ft.: IPE, ft.: t.: cTOR: ft.:	t:	369263.30 1026.00 .3000 44.900 11.380 3.000 97.919 78.000 175.919 1281127. .000150 .0155 10.867	
						3.3889	_

Problem 3-3 Maximum Compressible Fluid Flow

Calculate the maximum flow rate of natural gas through a ruptured exchanger tube assuming a complete break near the tube sheet as shown in Figure 3-3. The data are:

- exchanger tubes = 3/4 inch, Schedule 160
- internal diameter, d inch = 0.614
- tube length, Lst, ft = 20
- pipe friction factor for complete turbulence, f = 0.026
- compressibility factor, Z = 0.9
- gas temperature, $^{\circ}F = 100$
- molecular weight, $M_w = 19.5$
- pressure in exchanger tubes, $P_1(psig) = 1110 psig (1124.7 psia)$
- relief valve set pressure on the shell side, P₂(psig) = 400 psig (414.7 psia)
- ratio of specific heat capacities, k = 1.27
- fluid viscosity, μ , cP = 0.012
- resistance coefficient due to fittings and valves, K = 2.026

Solution

The computer program PROG33 determines the maximum flow rate for any compressible fluid, having a design specification that is $\Delta P >$ 10% of the upstream pressure or compressible fluids at high velocities. The program calculates the maximum fluid flow rate for a given pipe size, length of pipe, and the fluid's physical properties. Table 3-10 shows the input data and computer output for the maximum flow rate of natural gas in the 0.614-inch pipe. The computed maximum flow rate of natural gas through the 0.614-inch pipe is 8397 lb/hr. The Mach number at the pipe inlet is 0.2074. At critical condition, the Mach number is 0.8874. The critical pressure of the compressible fluid is 242.3 psia, with a sonic velocity of 1346.7 ft/sec. The compressible fluid flow pattern through the pipe is **SUBSONIC**.

Table 3-10 Input Data and Computer Output for Maximum Compressible Fluid in a Pipe Line

DATA33.	DAT			
2.026	0.9	0.026 100.0 1.27	19.5 0.012	
1124.7 ******* PIF STF MAX FLU PIL FLU FLU FLU RES RES TOTI INI INI OUT PRI	414.7 COMPF SE INTER AIGHT I UD DENS SE FRICT UD DENS SE FRICT UD TEMF UD TEMF UD TEMF SID VISC SISTANCE SISTANCE SISTANCE SISTANCE SISTANCE SISTANCE SISTANCE SISTANCE SISTANCE SISTANCE SISTANCE SUSTET FLUI	1.27 RESSIBLE INAL DIA ENGTH C UID FLC SITY, LE IION FAC RESSIBL SPECIFIC COSITY, COEFF. SCOEFF. SCALCEFF. SCALCEFF. SCALCEFF.	0.012 FLUID FLOW CALCULATIONS IN A ************************************	
FLU MAC MAC REY CRI FLU	JID SONI CH MUMBH CH NUMBH (NOLDS N (NOLDS N (TICAL H JID FLOW	IC VELOC ER AT IN ER AT CH NUMBER: PRESSURE VIS:	ITY, ft/sec.:	1346.739 .2074 .8874 86293830. 242.305 SUBSONIC

Problem 3-4 Compressible Fluid Pressure Drop

What is the overall pressure drop for natural gas flowing at 27,000 lb/hr through a 6-inch Schedule 40 tail pipe to a header during relieving condition as shown in Figure 3-3? Relief valve set pressure, $P_1 = 400$ psig, during relieving condition is

 $P_1 = P_1 + 10\%$ accumulation + 14.7

= 454.7 psia

The data are:

- pipe internal diameter, d inch = 6.065
- tail pipe length, ft = 10
- gas flow rate, lb/h = 27000
- gas viscosity, μ , cP = 0.012
- gas compressibility factor, Z=0.9
- gas temperature, $^{\circ}F = 100$
- molecular weight of gas, $M_w = 19.5$
- ratio of specific heat capacities k=1.27
- gas inlet pressure, psia, 454.7
- pipe roughness, $\varepsilon = 0.00015$ ft
- resistance coefficient due to fittings and valves, K, 2.013
- pipe line resistance: $1 \times 90^{\circ}$ LR elbows
 - $1 \times \text{entrance loss}$
 - $1 \times \text{exit loss}$
 - 4.026 inch pipe reduction

Solution

The computer program PROG34 determines the overall pressure drop for the 6-inch tail pipe having a relieving rate of 27,000 lb/hr. Table 3-11 illustrates both the input data and computer output. The Mach number at inlet condition is 0.0169, and at the critical condition is 0.8874. The critical pressure is 7.985 psia and the overall pressure drop is 0.213 psi. The compressible fluid flow pattern through the pipe is **SUBSONIC**.

Problem 3-5

Calculate the pressure drop of a 5 mile length in a 6-inch (Schedule 40, ID=6.065 inch) pipe, for a 5000 bpd rate of salt water ($\gamma_w = 1.07$), having a gas flowing at 6 mmscfd of 0.65 specific gravity at a temperature of 110°F.

Table 3-11 Input Data and Computer Output for Compressible Fluid Flow Pressure Drop

DATA34.DAT

6.065	10.0	27000.0	2.013
0.012	0.9	100.0	19.5
0.00015	1.27	454.7	

COMPRESSIBLE FLUID FLOW CALCULATIONS IN A P	
PIPE INTERNAL DIAMETER, inch:	6.065
STRAIGHT LENGTH OF PIPE, ft.:	10.000
FLUID FLOWRATE, 1b/h.:	27000.00
FLUID DENSITY, 1b/ft^3:	1.641
FLUID VISCOSITY, CP:	.0120
PIPE ROUGHNESS, ft.:	.000
FLUID COMPRESSIBILITY FACTOR:	.9000
FLUID TEMPERATURE, OF:	100.000
FLUID MOLECULAR WEIGHT, Mw:	19.500
RATIO OF SPECIFIC HEAT CAPACITIES, Cp/Cv:	1.270
RESISTANCE COEFF. DUE TO FRICTIONAL LOSS:	.301
RESISTANCE COEFF. DUE TO FITTINGS + VALVES:	2.013
TOTAL RESISTANCE COEFFICIENT:	2.314
INLET FLUID PRESSURE, psia:	454.700
OUTLET FLUID PRESSURE, psia:	454.487
PRESSURE DROP, psi:	.213
FLUID VELOCITY, ft/sec.:	22.766
FLUID SONIC VELOCITY, ft/sec.:	1346.739
MACH MUMBER AT INLET:	.0169
MACH NUMBER AT CRITICAL CONDITION:	.8874
DARCY FRICTION FACTOR:	.0152
REYNOLDS NUMBER:	2340890.
CRITICAL PRESSURE, psia:	7.985
FLUID VISCOSITY, CP:	.0120
FLUID FLOW IS:	SUBSONIC

Physical properties are:

	Liquid	Gas	
Flow rate W, lb/h	77956	12434	
Density ρ, lb/ft ³	66.7	2.98	
Viscosity µ, cP	1.0	0.02	
Surface tension σ, dyne/cm	70.0		

Pipe length 5 miles = 26,400 ft.

Solution

The computer program PROG35 calculates the flow regime and the pressure drops for liquid and vapor phases. From the two-phase flow modulus, the overall pressure drop for the 6-inch pipe is calculated. Table 3-12 gives the input data and computer output of the two-phase

Table 3-12Input Data and Computer Output for Two-PhaseFluid Flow in a Pipe Line

DATA35	DAT			
66.7	26400.0 70.0 0.00015	12434.0	1.0 0.02	
*****	TWO-PHAS		E LOSS CALCULATION	IN A PIPE LINE **********
PI	PE INTERNAL	L DIAMETER	. inch:	6.065
	UIVALENT LI			.000
	TUAL LENGTH			26400.000
	TAL LENGTH			26400.000
LI	QUID FLOWRA	ATE, lb/hr	.:	77956.000
LI	QUID REYNO	LDS NUMBER	:	81105.
LI	QUID FRICT	ION FACTOR	:	.0201
PR	ESSURE DROI	P OF LIQUI), psi/100ft.:	.0818
GA	S FLOW RATE	E, lb/hr.:		12434.000
GA	S REYNOLDS	NUMBER:		646814.
	S FRICTION			.0159
PR	ESSURE DROI	P OF GAS,]	psi/100ft.:	.0367
	OW REGIME 3			ANNULAR
			D PHASE FLOW MODULU	
VE	LOCITY OF 1	FLUID IN P	IPE, ft./sec.:	7.391
BA	KER PARAME	FER IN THE	LIQUID PHASE:	40.768
BA	KER PARAME	TER IN THE	GAS PHASE:	9495.209
TW	O-PHASE FLO	W MODULUS	:	10.0264
PR	ESSURE DRO	P OF TWO-P	HASE MIXTURE, psi/1	00ft: .3684
ov				XTURE, psi: 97.2478
IN	DEX	926. IS L	ESS THAN 10000 PIP	E EROSION IS UNLIKELY

flow. The Lockhart-Martinelli two-phase flow modulus is 1.49. The Baker parameter in the liquid phase is 40.8, and the Baker parameter in the gas phase is 9495.2. These results indicate that the flow regime is **ANNULAR** (and can be observed from the Baker's plot, Figure 3-5). The two-phase modulus is 10.03 and the pressure drop of the two-phase mixture per 100 ft of pipe is 0.368 psi/100ft. The overall pressure drop of the two-phase mixture is 97.25 psi. In addition, the pipe is unlikely to encounter any corrosion-erosion because the computed index of 926 is less than 10,000.

Problem 3-6

Calculate the Froude numbers and flow conditions for the 2-, 4- and 6-inch (Schedule 40) vertical pipes having the following liquid and vapor flow rates and densities.

WL = 6930 lb/h	WG = 1444 lb/h
$\rho_{\rm L} = 61.8 \ \rm lb/ft^3$	$\rho_{\rm G}=0.135~lb/ft^3$

Solution

The computer program PROG36 calculates the Froude numbers for the liquid and vapor phases. In addition, PROG36 will determine whether the pipe is self venting or whether pulsation flow is encountered. Table 3-13 shows the results for the 2-, 4-, and 6-inch (Schedule 40) pipes. Table 3-14 gives a typical input data and computer output for the 2-inch (Schedule 40) pipe.

Problem 3-7

Determine the pressure drop for the 4-, 6-, and 8-inch (Schedule 40) condensate headers under the following conditions:

flow = 10,000 lb/h steam condensate pressure = 114.7 psia header pressure = 14.7 psia

Pipe Internal Diameter: inch	2.067	4.026	6.065
Liquid Flow Rate: lb/h	6930	6930	6930
Liquid Density: lb/ft ³	61.80	61.80	61.80
Vapor Flow Rate: lb/h	1444	1444	1444
Vapor Density: lb/ft ³	0.135	0.135	0.135
Pipe Area: ft ²	0.023	0.088	0.201
Liquid Velocity: ft/s.	1.337	0.352	0.155
Vapor Velocity: ft/s.	127.504	33.609	14.810
Froude Number for Liquid Phase	0.5682	0.1073	0.0385
Froude Number for Vapor Phase	2.5332	0.4784	0.1718
	Flow is pulse and this may result in pipe vibration.	Line is self- venting. Therefore no vibration problems are expected.	Line is self- venting. Therefore no vibration problems are expected.

Table 3-13 Vapor-Liquid Two-Phase Downflow

Table 3-14 Input Data and Computer Output for Vapor-Liquid Two-Phase Vertical Downflow

DATA36.	TAC	
2.067 1444.0	6930.0 61.8 0.135	
******	VAPOR-LIQUID TWO-PHASE VERTICAL DOWNFLOW	****
	PIPE INTERNAL DIAMETER, inch:	2.067
	LIOUID FLOWRATE, 1b/h.:	6930.
	LIQUID DENSITY, 1b/ft^3:	61.800
	VAPOR FLOWRATE, 1b/h.:	1444.
	VAPOR DENSITY, 1b/ft^3:	.135
	PIPE AREA, ft^2:	.023
	LIQUID VELOCITY, ft/s:	1.337
	VAPOR VELOCITY, ft/s:	127.504
	FROUDE NUMBER FOR LIQUID PHASE:	.5682
	FROUDE NUMBER FOR VAPOR PHASE:	2.5332
	FLOW IS PULSE AND THIS MAY RESULT IN PIPE VIBRAT	ION

Solution

The computer program PROG37 evaluates the pressure drop of any given condensate header. The program also determines whether the velocity of the flashed condensate mixture would cause deterioration in the header line. Table 3-15 illustrates the results for the 4-, 6-, and 8-inch headers. Table 3-16 shows a typical input data and computer output for the 4-inch (Schedule 40) pipe. The computed results show that the 4-inch pipe gives the velocity of the flashed condensate mixture to be 7055 ft/min. This indicates a possible deterioration in the pipe. For the 6- and 8-inch pipes, the velocities are 3109 ft/min, and 1795 ft/min respectively, indicating that the condensate pipe lines will not deteriorate.

Problem 3-8

Calculate the pressure drop in a 60 ft length of 1 1/2 inch (Schedule 40, ID = 1.610 inch), pipe packed with catalyst pellets 1/4 inch in diameter when 104.4 lb/h of gas is passing through the bed. The temperature is constant along the length of pipe at 260°C. The void fraction is 45% and the properties of the gas are similar to those of air at this temperature. The entering pressure is 10 atm [29].

Solution

The computer program PROG38 calculates the pressure drop in a 1.610-inch I.D. packed bed for varying length of catalyst pellets. For air at 260°C and 10 atm, the physical properties are:

Table 3-15Line Sizes for Flashing Steam Condensate

Pipe Internal Diameter: inch.	4.026	6.065	7.981
Total flow of mixture in condensate heater: lb/h.	10000	10000	10000
Steam condensate pressure before flashing: psia.	114.7	114.7	114.7
Flashed condensate header pressure: psia.	14.7	14.7	14.7
Weight fraction of condensate flashed to vapor.	0.135	0.135	0.135
Flashed Steam flowrate: lb/h.	1346	1346	1346
Flashed condensate liquid flowrate: lb/h.	8654	8654	8654
Temperature of Flashed condensate: °F	212.36	212.36	212.36
Flashed Steam Density: lb/ft ³	0.036	0.036	0.036
Flashed condensate liquid density: lb/ft ³	59.780	59.780	59.780
Density of mixture: lb/ft ³	0.267	0.267	0.267
Friction Factor:	0.0163	0.0149	0.0141
Pressure drop of flashed condensate mixture: psi/100 ft.	1.937	0.228	0.055
Velocity of flashed condensate mixture: ft/min.	7055	3109	1795
	Velocity is greater than 5000 ft/min. Deteriora- tion of the line is possible	Velocity is less than 5000 ft/min. The condensate header line will not deteriorate	Velocity is less than 5000 ft/min. The condensate header will not deteriorate

Table 3-16 Input Data and Computer Output for Flashing Steam Condensate

DATA37.DAT 4.026 10000.0 114.7 14.7

LINE SIZING FOR FLASHING STEAM CONDENSATE	
***************************************	******
PIPE INTERNAL DIAMETER, inch:	4.026
TOTAL FLOW OF MIXTURE IN CONDENSATE HEADER, 1b/h:	10000.
STEAM CONDENSATE PRESSURE BEFORE FLASHING, psia:	114.700
FLASHED CONDENSATE HEADER PRESSURE, psia:	14.700
WEIGHT FRACTION OF CONDENSATE FLASHED TO VAPOR:	.135
FLASHED STEAM FLOWRATE, lb/h.:	1346.
FLASHED CONDENSATE LIQUID FLOWRATE, 1b/h.:	8654.
TEMPERATURE OF FLASHED CONDENSATE, oF:	212.360
FLASHED STEAM DENSITY, 1b/ft^3:	.036
FLASHED CONDENSATE LIQUID DENSITY, 1b/ft^3:	59.780
DENSITY OF MIXTURE, 1b/ft^3:	.267
FRICTION FACTOR:	.0163
PRESSURE DROP OF FLASHED, psi:	
CONDENSATE MIXTURE, psi/100ft:	1.937
VELOCITY OF FLASHED CONDENSATE MIXTURE, ft/min.:	7055.
FLASHED CONDENSATE MIXTURE VELOCITY IS GREATER THAN	5000 ft./min.
DETERIORATION OF THE LINE IS POSSIBLE	

Table 3-17Pressure Drop in a 1 1/2 Packed Bed

Particle length inch	Reynolds number, N _{Re}	Friction factor f _p	Pressure drop ΔP , lb/in ²
0.250	4170	1.7860	68.603
0.300	4415	1.7840	64.720
0.350	4609	1.7825	61.951
0.400	4765	1.7815	59.877
0.450	4895	1.7806	58.266

Table 3-18
Input Data and Computer Output for for
Pressure Drop in a Packed Bed

DATA38.	DAT				
0.25	60.0 0.25 0.0278				
		PRESSURE DROP IN A PACKED BED			
******	********				
		D BED DIAMETER, ft.:	.134		
		I OF PACKED BED, ft.:	60.000		
	FLUID	FLOWRATE THROUGH PACKED BED, 1b/h.:	104.4		
	PARTIC	CLE DIAMETER, inch.:	.250		
PARTICLE LENGTH, inch.: .250					
FRACTIONAL VOIDS IN PACKED BED: .450					
DENSITY OF FLUID, 1b/ft^3: .4130					
FLUID VISCOSITY, CP: .0278					
		-SECTIONAL AREA OF BED, ft^2:	.0141		
		FICIAL MASS VELOCITY, 1b/h.ft^2:	7402.9		
		OF PARTICLES, ft^2:	.2045E-02		
		E OF PARTICLES, ft^3:	.7102E-05		
		ACKED BED SURFACE AREA, ft^2/ft^3 BED:	158.400		
		FIVE PARTICLE DIAMETER, ft.:	.0208		
		REYNOLDS NUMBER:	4170.		
		ION FACTOR FOR INTERMEDIATE FLOW:	1.7860		
		PRESSURE DROP IN PACKED BED, 1b/in^2:			

density $\rho = 0.413 \text{ lb}_m/\text{ft}^3$ viscosity $\mu = 0.0278 \text{ cP}$

Table 3-17 gives the Reynolds number, friction factor, and pressure drop of catalyst pellets of 0.25 inch and at different particle length. Table 3-18 shows a typical input data and computer output with PL = 0.25 inch. The simulation exercise gives a pressure drop of 68.603 lb/in². The results show that the pressure drop in a packed bed depends on size and shape of the particles.

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SUBROUTINE PROG31

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С	*		TTINGS SUCH AS					*
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С	*		= PIPE CONTRA	ACTING DI	AMETER, ft	•		*
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С	*		= FLUID FLOW					*
С	*	W =	= GAS FLOWRAN	rE, lb/h.				*
с	*	G =	ACCELERATION	ON DUE TO	GRAVITY,	32.174 ft/sec	2^2	*
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С	*		E VALVE		4	300.0	0.10	*
с	*		L VALVE		5	500.0	0.15	*
С	*	PLUG	J VALVE		6	1000.0	0.25	*
С	*	GLOB	BE VALVE		7	1500.0	4.00	*
С	*	DIA	PHRAM, DAM TYP	PE VALVE	8	1000.0	2.00	*
с	*	BUTT	FERFLY VALVE		9	800.0	0.25	*
č	*		G CHECK VALVE		10	1500.0	1.50	*
č	*		THRU TEE (flag	naed/weld		150.0	0.05	*
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14 0.0 1.00 с с * PIPE EXIT ***** CHARACTER*30 INFILE, OUTFIL CHARACTER CHOICE DIMENSION CHOFIT (14, 4) LOGICAL FITT COMMON /FILES/ INFILE, OUTFIL COMMON/DSCR1/ PID, RLST, Q, W, DEN, VIS, Z, PR1 COMMON/DSCR2/ V, LEQ, LST, L, DELTAH, DELP, DELPT COMMON/DATA1/ PR, D, RE COMMON/DATA2/ ICOUNT COMMON/DATA3/ CHOFIT COMMON/DATA3/ CD, ED, OD, CHOICE COMMON/DATA5/ F, K, KOT COMMON/DATA6/ THETA1, THETA2, PI REAL L, LEQ, RLST, K, OD, KOT, KTOTAL С CALCULATE PIPE CROSS-SECTIONAL AREA, ft^2 KTOTAL = 0.0LEQ = 0.0 PR = 0.00015 G = 32.2PI = 3.1415927D = PID/12AREA = (PI*(D**2))/4.0CHECK WHETHER THE PIPE ROUGHNESS IS FOR CARBON STEEL С č (i.e. PR = 0.00015ft). IF (PR .NE. PR1) THEN PR = PR1ENDIF CHECK WHETHER FLUID FLOW RATE IS IN VOLUMETRIC OR MASS RATE. с IF (Q .GT. 0.0) THEN GO TO 11 ELSEIF (W .GT. 0.0) THEN GO TO 12 ENDIF с CALCULATE REYNOLDS NUMBER BASED ON VOLUMETRIC RATE 11 RE = (50.6*DEN*Q)/(PID*VIS)с CALCULATE FLUID VELOCITY ft/sec. V = (0.408*Q)/PID**2CALCULATE THE FLUID MASS FLOW RATE, 1b/hr С W = (V*(PID**2)*DEN)/.0509

GO TO 21 с CALCULATE REYNOLDS NUMBER BASED ON MASS FLOW RATE, 1b/hr. 12 RE = (6.31*W)/(PID*VIS)с CALCULATE FLUID VELOCITY, ft/sec. V = (0.0509 * W) / (DEN * (PID * * 2))С CALCULATE THE FLUID VOLUMETRIC FLOW RATE, gal/min. Q = W/(8.02 * DEN)С CALCULATE THE DARCY FRICTION FACTOR 21 IF (RE .GT. 2000.0) THEN GO TO 31 ELSE F1 = 64.0/REF = F1ENDIF GO TO 32 с CALCULATE THE FRICTION FACTOR USING THE CHEN'S EQUATION A1 = ((PR/D)/3.7+(6.7/RE)**0.9) B1 = -4.0*ALOGIO((PR/(3.7*D))-(5.02/RE)*ALOGIO(A1)) 31 С CALCULATE THE DARCY FRICTION FACTOR. F F = 4.0/B1**2С CHECK WHETHER THERE ARE FITTINGS ċ IF FIT IS TRUE THEN CALL THE PROGRAM SFIT FITT = .TRUE. IF (FITT) THEN 32 THEN CALL FITTINGS с CALL SFIT(K) GO TO 33 ELSE K = 0.0GO TO 34 ENDIF FITT = .FALSE.

CALCULATE THE TOTAL EQUIVALENT LENGTH , ft.

С

33 LEQ = LEQ + (K*D)/Fс CALCULATE THE TOTAL VELOCITY HEAD KTOTAL = KTOTAL + K + (F*RLST/D)CALCULATE TOTAL FITTING EQUIVALENT LENGTH PLUS С с STRAIGHT PIPE LENGTH, ft 34 L = RLST + LEOIF (RE .LT. 2000.0) THEN GO TO 35 ELSE ENDIF С CALCULATE THE FLUID PRESSURE DROP PER 100 ft. OF PIPE BASED ON THE Ċ MASS FLOW RATE, 1b/hr. IF (Q .GT. 0.0) THEN GO TO 44 ELSE DELP = (0.000336*F*(W**2))/((PID**5)*DEN) ENDIF с CALCULATE THE OVERALL PRESSURE DROP OF THE PIPE DELPT = (DELP*L)/100.0 + ((Z*DEN)/144.0)GO TO 55 CALCULATE THE FLUID PRESSURE DROP PER 100ft OF PIPE BASED ON THE С с VOLUMETRIC RATE. gal/min. 44 DELP = (0.0216*F*DEN*(Q**2))/(PID**5) с CALCULATE THE OVERALL PRESSURE DROP OF THE PIPE DELPT = (DELP*L)/100.0 + ((Z*DEN)/144.0) GO TO 55 с CALCULATE THE PRESSURE DROP FOR LAMINAR FLOW IF (Q .GT. 0.0) THEN 35 DELP = (0.0273 * VIS * Q) / (PID * * 4)С CALCULATE THE OVERALL PRESSURE DROP psi. DELPT = (DELP*L)/100.0 + ((Z*DEN)/144.0)С CALCULATE THE TOTAL HEAD LOSS, ft. DELTAH = (0.0393*VIS*L*Q)/(PID**4*DEN) + Z

GO TO 66 C C CALCULATE THE PRESSURE DROP BASED ON THE MASS FLOW RATE. THE PRESSURE DROP IN psi. PER 100ft OF PIPE ELSEIF (W .GT. 0.0) THEN DELP = (0.0034*VIS*W)/((PID**4)*DEN) С CALCULATE THE OVERALL PRESSURE DROP IN psi. DELPT = (DELP*L)/100.0 + ((Z*DEN)/144.0)С CALCULATE THE TOTAL HEAD LOSS, ft. $DELTAH = (0.0049 \times VIS \times L \times W) / ((PID \times 4) \times (DEN \times 2)) + Z$ ENDIF GO TO 66 С CALCULATE THE TOTAL HEAD LOSS ft. 55 DELTAH = (0.000483*F*L*(W**2))/((PID**5)*(DEN**2)) + Z 66 RETURN END С С * THIS PROGRAM USES THE TWO-K METHOD TO CALCULATE THE TOTAL C C C * THE VELOCITY HEAD FOR THE FITTINGS: * * CONSTANTS FOR THE TWO-K METHOD С SUBROUTINE SFIT (KALL) CHARACTER CHOICE DIMENSION CHOFIT (14, 4) COMMON/DATA1/ PR, D, RE COMMON/DATA3/ CHOFIT COMMON/DATA3/ CHOFIT COMMON/DATA4/ CD, ED, OD, CHOICE COMMON/DATA5/ F, K, KOT COMMON/DATA6/ THETA1, THETA2, PI COMMON/DSCR1/ PID, RLST, Q, W, DEN, VIS, Z, PR1 REAL KALL, KSUM, KLSUM, KEE1, KEE2, KEET, KTOTAL REAL KEX, KCT, KOD, KOT, OD, K CHOFIT (6, 3) = CHOFIT (1, 3)CHOFIT (8, 3) = CHOFIT (1, 3)CHOFIT (8, 3) = CHOFIT (1, 3)CHOFIT (9, 3) = CHOFIT (2, 3)CHOFIT (5, 3) = CHOFIT (3, 3)CHOFIT (10, 3) = CHOFIT (7, 3)CHOFIT (5, 4) = CHOFIT (3, 4)CHOFIT (9, 4) = CHOFIT (6, 4)

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KSUM = 0.0
           KLSUM = 0.0
           KEE1 = 0.0
KEE2 = 0.0
           KTOTAL = 0.0
           KEX = 0.0
           KCT = 0.0
           KOD = 0.0
            DO 10 I = 1, 12
           KSUM = KSUM + CHOFIT (I, 2)*CHOFIT (I, 3)
KLSUM = KLSUM + CHOFIT (I, 2)*CHOFIT (I, 4)
 10
          CONTINUE
           KTOTAL = KTOTAL + (KSUM/RE+(KLSUM*(1.0+(1.0/PID))))
KEE1=KEE1+(CHOFIT(13,2)*CHOFIT(13,3)+CHOFIT(14,2)*CHOFIT(14,3))
KEE2=KEE2+(CHOFIT(13,2)*CHOFIT(13,4)+CHOFIT(14,2)*CHOFIT(14,4))
           KEET = (KEE1/RE + KEE2)
           KALL = KTOTAL + KEET
           CALCULATE THE K VALUES OF THE SUDDEN CHANGES
С
с
           IN PIPE ENLARGEMENT/REDUCTION WITH TAPERED ANGLES OR WITHOUT TAPERED ANGLES.
с
           IF ( CHOICE .EQ. 'Y' .OR. CHOICE .EQ. 'y') THEN
           GO TO 20
           ELSEIF (CHOICE .EQ. 'N' .OR. CHOICE .EQ. 'n') THEN
           GO TO 75
           ENDIF
 20
          IF ( ED .EQ. 0.0) THEN
                    BETA1 = 0.0
           GO TO 30
           ELSE
           BETA1 = PID/ED
           ENDIF
                     IF (THETA1 .EQ. 0.0) THEN
                    GO TO 11
ELSEIF (THETA1 .LE. 45.0) THEN
CAL1 = 2.6*SIN((THETA1*PI)/(180.0*2.0))
                      GO TO 22
                     ELSEIF (THETA1 .GT, 45.0) THEN
                     ENDIF
 11
                     IF (RE .LE. 4000.0) THEN
                               BETA1 = 2.0*(1.0 - (BETA1)**4)
                               GO TO 30
                     ELSEIF (RE .GT. 4000.0) THEN
BETA1 = (1.0 + 0.8*F)*(1.0 - (BETA1**2))**2
                               GO TO 30
                     ENDIF
 22
                     IF (RE .LE. 4000.0) THEN
                               BETA1 = CAL1*2.0*(1.0 - (BETA1)**4)
```

		GO TO 30 ELSEIF (RE .GT. 4000.0) THEN BETA1 = CAL1*(1.0+0.8*F)*(1.0-(BETA1**2))**2 ENDIF
30	KEX = K	EX + BETAl
		IF (CD .EQ. 0.0) THEN BETA2 = 0.0 GO TO 40 ELSE BETA2 = PID/CD ENDIF
		IF (THETA2 .EQ. 0.0) THEN GO TO 35 ELSEIF (THETA2 .LT. 45.0) THEN CAL2 = 1.6*SIN((THETA2*PI)/(180.0*2.0))
		GO TO 45 ELSEIF (THETA2 .GT. 45.0 .OR. THETA2 .LT. 180.0) THEN CAL3 = SIN((THETA2*PI)/(180.0*2.0))
		CAL3 = SQRT(CAL3) GO TO 55 ENDIF
35		IF (RE .LE. 2500.0) THEN BETA2 = (1.2 + 160.0/RE)*((BETA2**4) - 1.0) GO TO 40 ELSEIF (RE .GT. 2500.0) THEN BETA2 = (0.6+0.48*F)*(BETA2**2)*((BETA2**2)-1.0) GO TO 40
		ENDIF
45		IF (RE .LE. 2500.0) THEN BETA2 = CAL2*(1.2+160.0/RE)*((BETA2**4)-1.0) GO TO 40 ELSEIF (RE .GT. 2500.0) THEN BETA2 = CAL2*(0.6+0.48*F)*(BETA2**2)*((BETA2**2)-1.0) GO TO 40 ENDIF
55		IF (RE .LE. 2500.0) THEN BETA2 = CAL3*(1.2+160.0/RE)*((BETA2**4)-1.0) GO TO 40 ELSEIF (RE .GT. 2500.0) THEN BETA2 = CAL3*(0.6+0.48*F)*(BETA2**2)*((BETA2**2)-1.0)
40		ENDIF KCT = KCT + BETA2
		IF (OD .EQ. 0.0) THEN BETA3 = 0.0 GO TO 41 ELSE BETA3 = OD/PID ENDIF
		IF (RE .LE. 2500.0) THEN VAL1 = (2.72+((BETA3**2)*(120.0/RE-1.0)))

	BETA3 = VAL1*(1.0-(BETA3**2))*((1.0/(BETA3**4))-1.0)
	GO TO 42 ELSEIF (RE .GT. 2500.0) THEN VAL2 = (2.72-((BETA3**2)*(4000.0/RE))) BETA3 = VAL2*(1.0-(BETA3**2))*((1.0/(BETA3**4))-1.0)
	ENDIF
42	KOD = KOD + BETA3
41	KALL = KALL + KEX + KCT + KOD + KOT
75	RETURN END
c c c	<pre>************************************</pre>

PROGRAM PROG32

c	****
С	
с	* THIS PROGRAM CALCULATES THE PRESSURE DROP IN A PIPE INCLUDING *
С	* PIPE FITTINGS SUCH AS VALVES AND ELBOWS. THE PROGRAM USES THE *
с	* CHEN'S METHOD TO DETERMINE THE FRICTION FACTOR - f. THE DARCY *
с	* FRICTION FACTOR IS 4FC (i.e. 4 X CHEN'S FRICTION FACTOR). *
с	* THE PROGRAM USES THE VELOCITY HEAD DUE TO PIPE FITTINGS+VALVES *
С	* TO CALCULATE THE EQUIVALENT PIPE LENGTH. THE TOTAL LENGTH OF *
с	* PIPE IS THE SUMMATION OF THE EQUIVALENT LENGTH AND THE STRAIGHT *
С	* PIPE LENGTH. FROM THE TOTAL LENGTH OF THE PIPE, THE OVERALL *
с	* PRESSURE DROP OF THE PIPE IS CALCULATED. *
c	***************************************
c	
č	*********************
č	* NOMENCLATURE *
č	* *
c	* *
с	
С	* PID = PIPE INTERNAL DIAMETER, inch. *
С	* ED = PIPE EXPANSION DIAMETER, ft. *
с	* CD = PIPE CONTRACTING DIAMETER, ft. *
с	* F = DARCY FRICTION FACTOR *
с	* L = TOTAL FITTING EQUIVALENT LENGTH PLUS *
с	* STRAIGHT PIPE LENGTH, ft. *
с	* LEQ = FITTINGS EQUIVALENT LENGTH, ft. *
с	* LST = STRAIGHT PIPE LENGTH, ft. *
с	* NF = NUMBER OF FITTINGS OF A GIVEN TYPE *
С	* RE = REYNOLDS NUMBER *
с	* $P = P1 \text{ OR } P2 \text{ (DEPENDING ON INPUT)}$.
c	* P1 = PIPE INLET PRESSURE, psia. *
c	* P2 = PIPE OUTLET PRESSURE, psia. *
c	* DELP = PRESSURE DROP psi/100 ft. *
č	* RHO = FLUID DENSITY, lb/ft^3. *
č	* Q = FLUID VOLUMETRIC FLOW RATE, gal/min. *
č	* W = FLUID MASS FLOWRATE, lb/h.
c	* G = ACCELERATION DUE TO GRAVITY, 32.174 ft/sec^2.
с	<pre>* PR = ABSOLUTE ROUGHNESS OF PIPE WALL, 0.00015ft. * * V = FLUID VELOCITY, ft/sec. *</pre>
С	
С	* VIS = VISCOSITY OF FLUID, cP. *
С	***************************************
	REAL RL, RLEQ, RLST, RK, KTOTAL
	COMMON/DATA1/ PID, RLST, Q, W, DEN, VIS, Z, PR1
	COMMON/DATA2/ V, RLEQ, KTOTAL, RK, RL, DELTAH, DELP, DELPT
	COMMON/DATA2/ V, REEQ, RIGIAL, RK, RE, DELTAR, DEEP, DEEPI COMMON/DATA3/ G, PR, D, AREA, RE, FF
	COMMON/DATAS/ G, PR, D, AREA, RE, FF
с	OPEN THE DATA FILE
C	OPEN THE DATA FILE
	OPEN (UNIT = 3, FILE = 'DATA32.DAT', STATUS = 'OLD', ERR = 18)
с	OPEN FILE FOR PRINTING THE RESULTS.
	OPEN (UNIT = 1, FILE = 'PRN')
FO	WRITE (1, 50)
50	FORMAT (///,\)

218

WRITE (1, 100) FORMAT (5X, 'PRESSURE DROP CALCULATION OF AN INCOMPRESSIBLE', \) 100 WRITE (1, 110) FORMAT (' FLUID IN A PIPE LINE',/1H , 78(1H*)) 110 с READ IN THE DATA IN THE DATA FILE. с READ THE PIPE INTERNAL DIAMERTER, inch: PID READ THE ACTUAL LENGTH OF PIPE, ft.: RLST 0 0 0 0 READ THE ACTUAL LENGTH OF FIFE, IC.: REST READ THE FLUID FLOW RATE (IF VOLUMETRIC FLOWRATE, gal/min.: Q IS GIVEN THEN THE MASS FLOW RATE, lb/hr.: W = 0.0, ALTERNATIVELY, IF THE MASS FLOW RATE, lb/hr.: W IS GIVEN, THEN THE VOLUMETRIC RATE, gal/min. Q = 0.0.00000 READ THE FLUID DENSITY, lb/ft^3 : DEN READ THE FLUID VISCOSITY, cP: VIS READ THE PIPE ELEVATION, ft.: Z (OTHERWISE Z = 0.0) READ THE VELOCITY HEAD DUE TO PIPE FITTINGS, :RK с READ THE PIPE ROUGHNESS, ft. PR1 (0.00015) READ (3, *, ERR=19) PID, RLST, Q, W READ (3, *, ERR=19) DEN, VIS, Z, RK READ (3, *, ERR=19) PR1 GO TO 5 18 WRITE (*, 120) 120 FORMAT (5X, 'DATA FILE DOES NOT EXIST') GO TO 999 19 WRITE (*, 130) FORMAT (5X, 'ERROR MESSAGE IN THE DATA VALUE') 130 GO TO 999 c c CALCULATE PIPE CROSS-SECTIONAL AREA, ft^2 5 PI = 3.1415927 G = 32.2D = PID/12AREA = (PI*(D**2))/4.0CHECK WHETHER THE PIPE ROUGHNESS IS FOR CARBON STEEL с с (i.e. PR=0.00015ft). IF (PR .NE. PR1) THEN PR = PR1 ENDIF с CHECK WHETHER FLUID FLOW RATE IS IN VOLUMETRIC OR MASS RATE.

IF (Q .GT. 0.0) THEN CALL VRATE CALL VMOUT ELSE CALL MRATE CALL VMOUT ENDIF CLOSE (UNIT = 3, STATUS = 'KEEP') CLOSE (UNIT = 1)999 STOP END с ****** c c c c THIS PROGRAM CALCULATES THE PRESSURE DROP OF INCOMPRESSIBLE FLUIDS. FOR A GIVEN VOLUMETRIC RATE. ********** SUBROUTINE VRATE REAL RL, RLEQ, RLST, RK, KTOTAL COMMON/DATA1/ PID, RLST, Q, W, DEN, VIS, Z, PR1 COMMON/DATA2/ V, RLEQ, KTOTAL, RK, RL, DELTAH, DELP, DELPT COMMON/DATA3/ G, PR, D, AREA, RE, FF RLEQ = 0.0KTOTAL = 0.0С CALCULATE THE REYNOLDS NUMBER RE = (50.6*DEN*Q)/(PID*VIS)С CALCULATE THE FLUID VELOCITY, ft/sec. $V = (0.408 \star 0) / PID \star \star 2$ с CALCULATE THE FLUID MASS RATE, 1b/hr. W = (V*PID**2*DEN)/0.0509 с DETERMINE WHETHER FLUID FLOW IS LAMINAR OR TURBULENT. IF (RE .GT. 2000.0) THEN IF FLUID FLOW IS TURBULENT, CALCULATE THE FRICTION FACTOR С с USING CHEN'S EQUATION. CALL CHEN ELSE с FLUID FLOW IS LAMINAR, CALCULATE THE FRICTION FACTOR.

FF = 64.0/RE ENDIF

C CALCULATE THE TOTAL EQUIVALENT LENGTH, ft.

CALL TLENGTH

C CALCULATE THE FLUID PRESSURE DROP PER 100ft. OF PIPE, psi/100ft. C CHECK WHETHER FLUID FLOW IS LAMINAR OR TURBULENT.

IF (RE .LT. 2000.0) THEN

DELP = (0.0273*VIS*Q)/(PID**4)

C CALCULATE THE OVERALL PRESSURE DROP, psi.

DELPT = (DELP*RL)/100.0+((Z*DEN)/144.0)

C CALCULATE THE TOTAL HEAD LOSS, ft.

DELTAH = (0.0393*VIS*RL*Q)/((PID**4)*DEN) + Z ELSE

- C IF FLUID FLOW IS TURBULENT C CALCULATE THE PRESSURE DROP PER 100ft. OF PIPE, psi/100ft.
- DELP = (0.0216 * FF * DEN * (Q * * 2)) / (PID * * 5)
- C CALCULATE THE OVERALL PRESSURE DROP, psi.

$$DELPT = (DELP*RL)/100.0 + ((Z*DEN)/144.0)$$

C CALCULATE THE OVERALL HEAD LOSS, ft.

```
DELTAH = (0.0311*FF*RL*(Q**2))/(PID**5) + Z
ENDIF
RETURN
END
```

SUBROUTINE MRATE REAL RL, RLEQ, RLST, RK, KTOTAL

COMMON/DATA1/ PID, RLST, Q, W, DEN, VIS, Z, PR1 COMMON/DATA2/ V, RLEQ, KTOTAL, RK, RL, DELTAH, DELP, DELPT COMMON/DATA3/ G, PR, D, AREA, RE, FF

 $\begin{array}{l} \text{RLEQ} = 0.0 \\ \text{KTOTAL} = 0.0 \end{array}$

C CALCULATE THE REYNOLDS NUMBER

RE = (6.31*W)/(PID*VIS)

с	CALCULATE THE FLUID VELOCITY, ft/sec.
	V = (0.0509*W)/(DEN*PID**2)
с	CALCULATE THE FLUID VOLUMETRIC FLOWRATE, gal/min.
	$Q = W/(8.02 \star DEN)$
с	DETERMINE WHETHER FLUID FLOW IS LAMINAR OR TURBULENT.
	IF (RE .GT. 2000.0) THEN
c c	IF FLUID FLOW IS TURBULENT, CALCULATE THE FRICTION FACTOR, USING CHEN'S EQUATION.
	CALL CHEN ELSE
С	IF FLUID FLOW IS LAMINAR, CALCULATE THE FRICTION FACTOR.
	FF = 64.0/RE ENDIF
с	CALCULATE THE TOTAL EQUIVALENT LENGTH, ft.
	CALL TLENGTH
с	CALCULATE THE FLUID PRESSURE DROP PER 100ft. OF PIPE, $psi/100ft$.
	IF (RE .LT. 2000.0) THEN
с	CALCULATE THE PRESSURE DROP FOR LAMINAR FLOW, psi/100ft.
	<pre>DELP = (0.0034*VIS*W)/((PID**4)*DEN)</pre>
с	CALCULATE THE OVERALL PRESSURE DROP FOR LAMINAR FLOW, psi.
	DELPT = (DELP*RL)/100.0 + ((Z*DEN)/144.0)
с	CALCULATE THE HEAD LOSS FOR LAMINAR FLOW.
	DELTAH = (0.0049*VIS*RL*W)/((PID**4)*(DEN**2)) + Z ELSE
с	CALCULATE THE PRESSURE DROP FOR TURBULENT FLOW psi/100ft.
	DELP = (0.000336*FF*(W**2))/((PID**5)*DEN)
с	CALCULATE THE OVERALL PRESSURE DROP, psi.
	DELPT = ((DELP*RL)/100.0) + ((Z*DEN)/144.0)
С	CALCULATE THE TOTAL HEAD LOSS, ft.
	DELTAH = (0.000483*FF*RL*(W**2))/((PID**5)*(DEN**2)) + Z ENDIF
	RETURN

END

с с с с с		**************************************
		SUBROUTINE VMOUT
		REAL RL, RLEQ, RLST, RK, KTOTAL
		COMMON/DATA1/ PID, RLST, Q, W, DEN, VIS, Z, PR1 COMMON/DATA2/ V, RLEQ, KTOTAL, RK, RL, DELTAH, DELP, DELPT COMMON/DATA3/ G, PR, D, AREA, RE, FF
140	* * *	<pre>WRITE (1,140) PID, W, Q, VIS FORMAT(5X,'PIPE INTERNAL DIAMETER, inch:', T60, F8.3,/, 5X,'MASS FLOW RATE, lb/h.:',T60, F10.2,/, 5X,'VOLUMETRIC FLOW RATE, gal/min.:',T60,F10.2,/, 5X,'FLUID VISCOSITY, cP:',T60,F8.4)</pre>
150	* * *	<pre>WRITE (1, 150) DEN, V, RK, RLEQ FORMAT(5X,'FLUID DENSITY, lb/ft^3:', T60, F8.3,/, 5X,'FLUID VELOCITY, ft/sec.:',T60, F8.3,/, 5X,'VELOCITY HEAD LOSS DUE TO FITTINGS:',T60, F8.3,/, 5X,'EQUIVALENT LENGTH OF PIPE, ft.:',T60, F8.3)</pre>
160	* * *	<pre>WRITE (1, 160) RLST, RL, RE, PR FORMAT(5X,'ACTUAL LENGTH OF PIPE, ft.:',T60, F8.3,/, 5X,'TOTAL LENGTH OF PIPE, ft.:', T60, F8.3,/, 5X,'REYNOLDS NUMBER:', T60, F10.0,/, 5X,'PIPE ROUGHNESS, ft.:', T60, F8.6)</pre>
170	* * *	<pre>WRITE (1, 170) FF, DELTAH, DELP, DELPT FORMAT(5X,'DARCY FRICTION FACTOR:', T60, F8.4,/, 5X,'EXCESS HEAD LOSS, ft.:', T60, F8.3,/, 5X,'PIPE PRESSURE DROP /100ft., psi/100ft:', T60, F8.4,/, 5X,'OVERALL PRESSURE DROP OF PIPE, psi:', T60, F8.4)</pre>
180		WRITE (1, 180) FORMAT (78(1H-))
с		FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
		WRITE (1, *) CHAR(12)
		RETURN END
с с с с		**************************************

		SUBROUTINE CHEN
		COMMON/DATA3/ G, PR, D, AREA, RE, FF

```
A1 = ((PR/D)/3.7+(6.7/RE)**0.9)
B1 = -4.0*ALOG10((PR/(3.7*D))-(5.02/RE)*ALOG10(A1))
         FF = 4.0/B1**2
         RETURN
         END
C
C
C
C
C
         THIS PROGRAM CALCULATES THE TOTAL LENGTH OF PIPE INCLUDING
         THE EQUIVALENT LENGTH.
         SUBROUTINE TLENGTH
        REAL RL, RLEQ, RLST, RK, KTOTAL
        COMMON/DATA1/ PID, RLST, Q, W, DEN, VIS, Z, PR1
COMMON/DATA2/ V, RLEQ, KTOTAL, RK, RL, DELTAH, DELP, DELPT
COMMON/DATA3/ G, PR, D, AREA, RE, FF
        \begin{array}{l} \text{RLEQ} \ \simeq \ 0.0 \\ \text{KTOTAL} \ = \ 0.0 \end{array}
с
        CALCULATE THE TOTAL EQUIVALENT LENGTH, ft.
         RLEQ = RLEQ+(RK*D)/FF
         CALCULATE THE TOTAL FITTING EQUIVALENT LENGTH PLUS STRAIGHT PIPE
С
č
        LENGTH, ft.
        RL = RLST + RLEQ
с
        CALCULATE THE VELOCITY HEAD, KTOTAL
        KTOTAL = KTOTAL + RK + (FF*RLST/D)
        RETURN
         END
```

PROGRAM PROG33

* THIS PROGRAM CALCULATES THE MAXIMUM FLUID RATE OF A * * COMPRESSIBLE FLUID FOR A GIVEN PIPE SIZE, PRESSURE DROP AND * * FLUID FLOW CHARACTERISTICS. * NOMENCLATURE * * * * * * * PIPE CROSS-SECTIONAL AREA, ft^2. * Α -× d = PIPE INTERNAL DIAMETER, inch. PIPE INTERNAL DIAMETER, ft. * × D = * * PFF = PIPE FRICTION FACTOR * ACCELERATION DUE TO GRAVITY, 32.2 ft/sec^2. q = ٠ * CONVERSION FACTOR (32.174 (lbm/lbf)(ft/s^2) g * ć * KF = RESISTANCE COEFFICIENT DUE TO PIPE FITTINGS+VALVES KP = RESISTANCE COEFFICIENT DUE TO FIPE FRICTION KTOTAL = TOTAL RESISTANCE COEFFICIENT × * * * = LENGTH OF STRAIGHT PIPE, ft. = MOLECULAR WEIGHT OF FLUID (lb/lb mol.) × LST * * MW * = INLET FLUID PRESSURE, psia. * P1 * OUTLET FLUID PRESSURE, psia.
 PRESSURE DROP (PRESSURE DIFFERENCE BETWEEN TWO **P**2 * DELP POINTS), psi. = MOLAR GAS CONSTANT (10.73 psia.ft^3/R.lb mole) * * × R × = 1544 lbf.ft/oR. lb mole. = MOLECULAR WEIGHT OF GAS/MOLECULAR WEIGHT OF AIR * ٠ SG × * = FLUID TEMPERATURE: oF. = FLUID VELOCITY, ft/sec. * т * * VG * * vs FLUID SONIC VELOCITY, ft/sec. = * = FLUID DENSITY, lb/ft^3. = FLUID VISCOSITY, cP. × RHO × * VIS * -FLUID COMPRESSIBILITY FACTOR (Z=1). z *********** ********************

CHARACTER*10 REGIME(3), REG(3)

Supersonic).

COMMON/DATA1/ D, RE COMMON/DATA2/ RHO, DELP COMMON/DATA3/ VG, VS, MACH, MACR, PC COMMON/DATA4/ KP, KTOTAL COMMON/DATA5/ REG, REGIME, IR

REAL ID, LST, KF, KP, MW, MACH, MACR, KTOTAL

C C

DATA REG/ 'SUBSONIC ', 'SONIC ', 'SUPERSONIC'/

FLOW PATTERNS OF COMPRESSIBLE FLUID FLOW IN A PIPE (Subsonic, Sonic

с OPEN THE DATA FILE. OPEN (UNIT = 3, FILE = 'DATA33.DAT', STATUS = 'OLD', ERR = 18) OPEN FILE FOR PRINTING THE RESULTS. С OPEN (UNIT = 1, FILE = 'PRN') WRITE (1, 100) 100 FORMAT (///,10X,'COMPRESSIBLE FLUID FLOW CALCULATIONS IN A', \) WRITE (1, 110) FORMAT (' PIPE LINE',/78(1H*)) 110 с READ IN THE DATA READ THE INTERNAL PIPE DIAMETER, inch.: ID READ THE STRAIGHT LENGTH OF PIPE, ft.: LST С С С READ THE PIPE FRICTION FACTOR: PFF READ THE RESISTANCE COEFFICIENT DUE TO PIPE FITTINGS + VALVES: KF READ THE FLUID COMPRESSIBILITY FACTOR: (Z=1.0) С С READ THE FLUID TEMPERATURE, OF: T READ THE FLUID MOLECULAR WEIGHT: MW 00000 READ THE FLUID INLET PRESSURE, psia: P1 READ THE FLUID OUTLET PRESSURE, psia: P2 READ THE RATIO OF SPECIFIC HEAT CAPACITIES, Cp/Cv: RATIOK с READ THE FLUID VISCOSITY, CP: VIS. READ (3, *, ERR=19) ID, LST, PFF READ (3, *, ERR=19) KF, Z, T, MW READ (3, *, ERR=19) P1, P2, RATIOK, VIS GO TO 20 18 WRITE (*, 120) FORMAT (5X, 'DATA FILE DOES NOT EXIST') 120 GO TO 999 WRITE (*, 130) FORMAT (5X, 'ERROR MESSAGE IN THE DATA VALUE') 19 130 GO TO 999 THIS CALLS FOR THE MAXIMUM FLUID FLOWRATE CALCULATION С 20 CALL CALC1 (ID, LST, PFF, KF, Z, T, MW, P1, P2, RATIOK, VIS, G) с THIS CALLS FOR THE RESULTS TO BE PRINTED. CALL OUT1 (ID, LST, PFF, KF, Z, T, MW, G, P1, P2, RATIOK, VIS) CLOSE (3, STATUS = 'KEEP') CLOSE (1) 999 STOP END

00000 THIS PROGRAM CALCULATES THE MAXIMUM COMPRESSIBLE FLUID RATE FOR A GIVEN PRESSURE LOSS AND FLUID SYSTEM CHARACTERISTICS. SUBROUTINE CALC1 (ID, LST, PFF, KF, Z, T, MW, P1, P2, RATIOK, * VIS, G) CHARACTER*10, REGIME(3), REG(3) COMMON/DATA1/ D, RE COMMON/DATA2/ RHO, DELP COMMON/DATA3/ VG, VS, MACH, MACR, PC COMMON/DATA4/ KP, KTOTAL COMMON/DATA5/ REG, REGIME, IR REAL ID, LST, KF, KP, KTOTAL, MW, MACH, MACR T = T + 460.0PI = 3.1415927 С CALCULATE THE PIPE AREA, ft^2. D = ID/12.0AREA = (PI*D*D)/4.0с CALCULATE THE INLET FLUID DENSITY, 1b/ft^3. RHO = (P1*MW)/(10.72*T*Z)с CALCULATE THE PRESSURE DIFFERENCE, psi. DELP = P1 - P2с CALCULATE THE RESISTANCE COEFFICIENT, KP DUE TO PIPE FRICTION. KP = (PFF*LST)/DCALCULATE THE TOTAL RESISTANCE COEFFICIENT. с KTOTAL = KF + KPCALCULATE THE MAXIMUM FLUID RATE FOR THE INLET AND OUTLET C C PRESSURES, psia. C1 = 1335.6*(ID**2)C2 = SQRT(RHO/(KTOTAL+(2.0*ALOG(P1/P2))))*SQRT((P1**2-P2**2)/P1) $G = C1 \star C2$ Ç CALCULATE THE FLUID'S REYNOLDS NUMBER AT THE MAXIMUM FLOW RE = (6.31*G)/(D*VIS)с CALCULATE THE FLUID VELOCITY, ft/sec. VG = (0.0509*G)/(RHO*ID**2)с CALCULATE THE SONIC VELOCITY, ft/sec. VS = 223.0*SQRT((RATIOK*T)/MW)

с CALCULATE THE INLET MACH NUMBER MACH = VG/VSCALCULATE THE EXIT MACH NUMBER FOR ISOTHERMAL FLOW С С i.e. THE MACH NUMBER AT CRITICAL CONDITION. MACR = 1.0/SQRT(RATIOK) CHECK WHETHER FLUID FLOW IS SUBSONIC, SONIC OR SUPERSONIC С IF (MACH .LT, MACR) THEN TR = 1REGIME(IR) = REG(IR) GO TO 10 ELSEIF (MACH .EQ. MACR) THEN IR = 2REGIME(IR) = REG(2)GO TO 10 ELSEIF (MACH .GT. MACR) THEN IR = 3REGIME(IR) = REG(3)ENDIF С CALCULATE THE CRITICAL PRESSURE AT SONIC CONDITION с USING CROCKER'S EQUATION, psia. SG = MW/29.010 R = 1544.0/(29.0*SG)PC = (G/(11400.0*ID**2))*SQRT((R*T)/(RATIOK*(RATIOK+1.0))) T = T - 460.0RETURN END С č THIS PROGRAM PRINTS THE RESULTS OF THE MAXIMUM COMPRESSIBLE c FLUID FLOW ONTO A PRINTER. SUBROUTINE OUT1 (ID, LST, PFF, KF, Z, T, MW, G, P1, P2, * RATIOK, VIS) CHARACTER*10, REGIME(3), REG(3) с COMMON/DATA1/ D, RE COMMON/DATA2/ RHO, DELP COMMON/DATA3/ VG, VS, MACH, MACR, PC COMMON/DATA4/ KP, KTOTAL COMMON/DATA5/ REG, REGIME, IR REAL ID, LST, KF, KP, KTOTAL, MW, MACH, MACR WRITE (1, 140) ID, LST, G, RHO, PFF, Z, T, MW, RATIOK, VIS FORMAT (5X,'PIPE INTERNAL DIAMETER, inch:', T60, F8.3,/, 5X,'STRAIGHT LENGTH OF PIPE, ft.:', T60, F8.3,/, 5X,'MAXIMUM FLUID FLOWRATE, lb/h.:', T60, F10.2,/, 140 1 2

3 4 5 6 7 8 9 *	5X,'FLUID DENSITY, lb/ft^3:', T60, F8.3,/, 5X,'PIPE FRICTION FACTOR:', T60, F8.4,/, 5X,'FLUID COMPRESSIBILITY FACTOR:', T60, F8.4,/, 5X,'FLUID TEMPERATURE, oF:', T60, F8.3,/, 5X,'FLUID MOLECULAR WEIGHT, Mw:', T60, F8.3,/, 5X,'RATIO OF SPECIFIC HEAT CAPACITIES, CP/Cv:',T60, F8.3,/, 5X,'FLUID VISCOSITY, CP:', T60, F8.4)
150 1 2	<pre>WRITE (1, 150) KP, KF, KTOTAL FORMAT (5X, 'RESISTANCE COEFF. DUE TO FRICTIONAL LOSS:',T60, F8.3,/,5X,'RESISTANCE COEFF. DUE TO FITTINGS + VALVES:', T60, F8.3,/,5X,'TOTAL RESISTANCE COEFFICIENT:',T60,F8.3)</pre>
160 1 2	<pre>WRITE (1,160) P1, P2, DELP FORMAT (5X,'INLET FLUID PRESSURE, psia:', T60, F9.3,/, 5X,'OUTLET FLUID PRESSURE, psia:', T60, F9.3,/, 5X,'PRESSURE DROP, psi:', T60, F9.3)</pre>
170 1 2 3	WRITE (1, 170) VG, VS, MACH, MACR FORMAT (5X,'FLUID VELOCITY, ft/sec.:', T60, F9.3,/, 5X,'FLUID SONIC VELOCITY, ft/sec.:', T60, F9.3,/, 5X,'MACH MUMBER AT INLET:', T60, F8.4,/, 5X,'MACH NUMBER AT CRITICAL CONDITION:', T60, F8.4)
180 1	WRITE (1, 180) RE, PC FORMAT (5X,'REYNOLDS NUMBER:', T60, F10.0/, 5X,'CRITICAL PRESSURE, psia:', T60, F9.3)
190	WRITE (1, 190) REGIME(IR) FORMAT (5X,'FLUID FLOW IS:', T60, A10)
200	WRITE (1, 200) Format (78(1H-))
с	FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR(12)
	RETURN END

* THIS PROGRAM CALCULATES THE PRESSURE DROP OF A × * COMPRESSIBLE FLUID FOR A GIVEN FLUID FLOW RATE, PIPE SIZE ٠ * AND FLUID FLOW CHARACTERISTICS. × NOMENCLATURE × * ------× * × PIPE CROSS-SECTIONAL AREA, ft^2. * × d PIPE INTERNAL DIAMETER, inch. * PIPE INTERNAL DIAMETER, ft. * D * DARCY FRICTION FACTOR FD * ACCELERATION DUE TO GRAVITY, 32.2 ft/sec^2. q * CONVERSION FACTOR (32.174 (lbm/lbf)(ft/s^2) q с * RESISTANCE COEFFICIENT DUE TO PIPE FITTINGS+VALVES RESISTANCE COEFFICIENT DUE TO PIPE FRICTION * KF = * KP = × = × KTOTAL TOTAL RESISTANCE COEFFICIENT × * LST z: LENGTH OF STRAIGHT PIPE, ft. MOLECULAR WEIGHT OF FLUID (1b/1b mol.) * MW = INLET FLUID PRESSURE, psia. OUTLET FLUID PRESSURE, psia. PRESSURE DROP (PRESSURE DIFFERENCE BETWEEN TWO * **P**1 * P2 * DELP = * POINTS), psi. PIPE ROUGHNESS (i.e. PR=0.00015ft.) * PR = MOLAR GAS CONSTANT (10.73 psia.ft^3/R.1b mole) * R = * MOLECULAR WEIGHT OF GAS/MOLECULAR WEIGHT OF AIR * ÷ * * SG = * * FLUID TEMPERATURE: DEG.F т FLUID VELOCITY, ft/sec. FLUID SONIC VELOCITY, ft/sec. * VG = * vs Ŧ FLUID DENSITY, 1b/ft^3. FLUID VISCOSITY, CP * RHO = * VIS = * FLUID COMPRESSIBILITY FACTOR (Z=1) Z = ** ************ *****

CHARACTER*10 REGIME(3), REG(3)

COMMON/DATA1/ D, RE, FD COMMON/DATA2/ RHO, DELP COMMON/DATA3/ VG, VS, MACH, MACR, PC COMMON/DATA4/ KF, KP, KTOTAL COMMON/DATA5/ REG, REGIME, IR

REAL ID, LST, KF, KP, MW, MACH, MACR, KTOTAL

C FLOW PATTERNS IN COMPRESSIBLE FLOW IN A PIPE (Subsonic, Sonic C Supersonic).

DATA REG/'SUBSONIC ', 'SONIC ', 'SUPERSONIC'/

с OPEN THE DATA FILE. OPEN (UNIT = 3, FILE = 'DATA34.DAT', STATUS = 'OLD', ERR = 18) OPEN FILE FOR PRINTING THE RESULTS. С OPEN (UNIT = 1, FILE = 'PRN') WRITE (1, 100) FORMAT (///,10X, 'COMPRESSIBLE FLUID FLOW CALCULATIONS IN A'. \) 100 WRITE (1, 110) FORMAT (' PIPE LINE',/78(1H*)) 110 с READ IN THE DATA с READ THE PIPE INSIDE DIAMETER, inch: ID READ THE STRAIGHT LENGTH OF PIPE, ft: LST С READ THE FLUID FLOW RATE, 1b/hr.: W READ THE RESISTANCE COEFF. DUE TO PIPE FITTINGS (e.g. BENDS, 0000 VALVES, ENTRANCE AND EXIT LOSES): KF READ THE FLUID VISCOSITY, CP: VIS READ THE FLUID COMPRESSIBILITY FACTOR: Z READ THE FLUID TEMPERATURE, OF: T с с READ THE FLUID MOLECULAR WEIGHT: MW C C C C READ THE PIPE ROUGHNESS, ft: PR1 READ THE RATIO OF SPECIFIC HEAT CAPACITIES, Cp/Cv: RATIOK č READ THE FLUID INLET PRESSURE, psia: P1 READ (3, *, ERR=19) ID, LST, W, KF READ (3, *, ERR=19) VIS, Z, T, MW READ (3, *, ERR=19) PR1, RATIOK, P1 GO TO 20 WRITE (*, 120) FORMAT (5X, 'DATA FILE DOES NOT EXIST') 18 120 GO TO 999 WRITE (*, 130) FORMAT (5X, 'ERROR MESSAGE IN THE DATA VALUE') GO TO 999 19 130 с THIS CALLS FOR THE PRESSURE DROP CALCULATION 20 CALL CALC2 (ID, LST, W, VIS, Z, T, MW, PR1, RATIOK, P1, P2) с THIS CALLS FOR THE RESULTS TO BE PRINTED. CALL OUT2 (ID, LST, W, Z, T, MW, RATIOK, VIS, PR1, P1, P2) CLOSE (3, STATUS = 'KEEP') CLOSE (1) STOP 999 END

с

CCCC THIS PROGRAM CALCULATES THE PRESSURE DROP OF A COMPRESSIBLE FLUID FOR A GIVEN FLOW RATE AND FLUID SYSTEM CHARACTERISTICS. ******* ******* ******** SUBROUTINE CALC2 (ID, LST, W, VIS, Z, T, MW, PR1, RATIOK, * P1, P2) CHARACTER*10, REGIME(3), REG(3) COMMON/DATA1/ D, RE, FD COMMON/DATA2/ RHO, DELP COMMON/DATA3/ VG, VS, MACH, MACR, PC COMMON/DATA4/ KF, KP, KTOTAL COMMON/DATA5/ REG, REGIME, IR REAL ID, LST, KF, KP, KTOTAL, MW, MACH, MACR PR = 0.00015T = T + 460.0PI = 3.1415927 с CALCULATE THE PIPE AREA, ft^2. D = ID/12.0AREA = (PI*D*D)/4.0с CALCULATE THE INLET FLUID DENSITY, 1b/ft^3. RHO = (P1*MW)/(10.72*T*Z)CHECK WHETHER THE PIPE ROUGHNESS IS FOR CARBON STEEL c c i.e. PR = 0.00015ftIF (PR .NE. PR1) THEN PR = PR1ENDIF С CALCULATE THE FLUID'S REYNOLDS NUMBER RE = (6.31*W)/(ID*VIS)IF (RE .GT. 2000.0) THEN A1 = ((PR/D)/3.7+(6.7/RE)**0.9) $B1 = -4.0 \times ALOG10((PR/(3.7 \times D)) - (5.02/RE) \times ALOG10(A1))$ FD = 4.0/B1**2ELSE FD = 64.0/REENDIF с CALCULATE THE RESISTANCE COEFFICIENT DUE TO PIPE FRICTION KP = (FD*LST)/DCALCULATE THE TOTAL RESISTANCE COEFFICIENT С с (i.e. FRICTIONAL LOSS + PIPE FLTTINGS) KTOTAL = KP + KFCALCULATE THE OUTLET PRESSURE OF THE FLUID, psia. С

```
с
        C1 = 1335.6*(ID**2)
        C2 = (W**2*P1*KTOTAL)/(C1**2*RHO)
        P2 = SQRT(P1 * * 2 - C2)
        CALCULATE THE PRESSURE DROP, psi.
С
        DELP = P1 - P2
        CALCULATE THE FLUID VELOCITY, ft/sec.
С
ċ
        VG = (0.0509 * W) / (RHO * ID * * 2)
с
        CALCULATE THE SONIC VELOCITY, ft/sec.
C
        VS = 223.0*SQRT((RATIOK*T)/MW)
        CALCULATE THE INLET MACH NUMBER
С
        MACH = VG/VS
        CALCULATE THE EXIT MACH NUMBER FOR ISOTHERMAL FLOW
C
C
        i.e. THE MACH NUMBER AT CRITICAL (CHOKING) CONDITION.
        MACR = 1.0/SQRT(RATIOK)
        CHECK WHETHER FLUID FLOW IS SUBSONIC, SONIC OR SUPERSONIC
с
        IF (MACH .LT. MACR) THEN
                TR = 1
                REGIME(IR)=REG(IR)
        GO TO 10
                    ELSEIF (MACH .EQ. MACR) THEN
                    IR = 2
                    REGIME(IR)=REG(2)
                    GO TO 10
                        ELSEIF (MACH .GT. MACR) THEN
                        IR = 3
                        REGIME(IR)=REG(3)
        ENDIF
        CALCULATE THE CRITICAL PRESSURE AT SONIC CONDITION USING CROCKER'S EQUATION, psia.
C
C
        SG = MW/29.0
10
        R = 1544.0/(29.0*SG)
        PC = (W/(11400.0*ID**2))*SQRT((R*T)/(RATIOK*(RATIOK+1.0)))
        T = T - 460.0
        RETURN
        END
        *********************************
                                                    ****************
C
C
C
        THIS PROGRAM PRINTS THE RESULTS OF THE MAXIMUM COMPRESSIBLE
        FLUID FLOW ONTO A PRINTER.
С
        *****************
                            SUBROUTINE OUT2 (ID, LST, W, Z, T, MW, RATIOK, VIS, PR1, P1, P2)
```

233

CHARACTER*10, REGIME(3), REG(3) COMMON/DATA1/ D, RE, FD COMMON/DATA2/ RHO, DELP COMMON/DATA3/ VG, VS, MACH, MACR, PC COMMON/DATA4/ KF, KF, KTOTAL COMMON/DATA5/ REG, REGIME, IR REAL ID, LST, MW, KF, KP, KTOTAL, MACH, MACR WRITE (1, 140) ID, LST, W, RHO, VIS, PR1, Z, T, MW, RATIOK FORMAT (5x,'PIPE INTERNAL DIAMETER, inch:', T60, F8.3,/, 5x,'STRAIGHT LENGTH OF PIPE, ft.:', T60, F8.3,/, 5x,'FLUID FLOWRATE, 1b/h.:', T60, F10.2,/, 5x,'FLUID DENSITY, 1b/ft^3:', T60, F8.3,/, 5x,'FLUID VISCOSITY, CP:', T60, F8.4,/, 5x,'PIPE ROUGHNESS, ft.:', T60, F8.3,/, 5x/FLUID CONDERSIBILITY FACTOR:', T60, F8.4,/, 140 1 2 3 4 5 5X, 'FLUID COMPRESSIBILITY FACTOR:', T60, F8.3,/, 5X,'FLUID COMPRESSIBILITY FACTOR:', T60, F8.4,/, 5X,'FLUID TEMPERATURE, oF:', T60, F8.3,/, 5X,'FLUID MOLECULAR WEIGHT, Mw:', T60, F8.3,/, 5X,'RATIO OF SPECIFIC HEAT CAPACITIES, Cp/Cv:',T60,F8.3) 6 7 8 9 WRITE (1, 150) KP, KF, KTOTAL FORMAT (5x,'RESISTANCE COEFF. DUE TO FRICTIONAL LOSS:',T60, F8.3,/,5x,'RESISTANCE COEFF. DUE TO FITTINGS + VALVES:', T60, F8.3,/,5x,'TOTAL RESISTANCE COEFFICIENT:', T60, F8.3) 150 1 WRITE (1, 160) P1, P2, DELP FORMAT (5X,'INLET FLUID PRESSURE, psia:', T60, F9.3,/, 5X,'OUTLET FLUID PRESSURE, psia:', T60, F9.3,/, 5X,'PRESSURE DROP, psi:', T60, F9.3) 160 1 2 WRITE (1, 170) VG, VS, MACH, MACR FORMAT (5X,'FLUID VELOCITY, ft/sec.:', T60, F9.3,/, 170 5X,'FLUID SONIC VELOCITY, ft/sec.:', T60, F3.3,', 5X,'FLUID SONIC VELOCITY, ft/sec.:', T60, F9.3,/, 5X,'MACH MUMBER AT INLET:', T60, F8.4,', 5X,'MACH NUMBER AT CRITICAL CONDITION:', T60, F8.4) 1 2 3 WRITE (1, 180) FD, RE FORMAT (5X,'DARCY FRICTION FACTOR:',T60, F8.4,/, 5X,'REYNOLDS NUMBER:', T60, F10.0) 180 WRITE (1, 190) PC, VIS FORMAT (5X, 'CRITICAL PRESSURE, psia:', T60, F9.3,/, 5X,'FLUID VISCOSITY, CP:', T60, F8.4) 190 WRITE (1, 200) REGIME(IR) 200 FORMAT (5X, 'FLUID FLOW IS:', T60, A10) WRITE (1, 210) FORMAT (78(1H-)) 210 FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE. c WRITE (1, *) CHAR(12) RETURN END

PROGRAM PROG35

		GRAM USES THE GENERAL METHOD OF LOCKHART AND MARTINELLI WITH OVID BAKER'S TWO-PHASE FLOW CORRELATIONS TO DETERMIN
*THE T	WO-F	PHASE PRESSURE DROP IN PROCESS PIPE LINES. THE PROGRAM
*ASSUM	ES I	HAT THE TWO-PHASE FLOW IS ISOTHERMAL AND TURBULENT IN BO
*LIQUI	D AN	ID VAPOR PHASES, AND THAT THE PRESSURE LOSS IS NOT MORE
*THAN	10 F	PER CENT OF THE ABSOLUTE UPSTREAM PRESSURE.
*THE P	ROGF	AM USES THE EXPLICIT EQUATIONS FOR FRICTION FACTOR
*DEVEL	OPEC	BY CHEN. IN ADDITION, THE PROGRAM DETERMINES WHETHER
		ND CORROSION IN THE PIPE MAY OCCUR.
*****	****	***************************************
*****	****	**************************************
*		NOMENCLATORE
*		
* RE	=	REYNOLDS NUMBER
* Wx		LIQUID OR GAS FLOW, 1b/h.
* ID		PIPE INSIDE DIAMETER, inch.
* VIS		VISCOSITY OF LIQUID OR GAS, CP.
* FF		FRICTION FACTOR
		PRESSURE DROP OF LIQUID OR GAS IF FLOWING IN THE PIPE
*	-	ALONE, psi/100 ft.
* DEN	=	DENSITY OF LIQUID OR GAS, 1b/ft^3.
* PR		PIPE ROUGHNESS, ft.
* VEL		VELOCITY OF FLUID IN PIPE, ft/s.
* DENG		GAS DENSITY, 1b/ft^3.
* DENL		LIQUID DENSITY, 1b/ft^3.
* WG	=	GAS FLOW, 1b/h.
* WL	=	LIQUID FLOW, 1b/h.
* A	=	CROSS-SECTIONAL AREA OF PIPE, ft^2.
		VISCOSITY OF LIQUID, CP.
* VISG		VISCOSITY OF VAPOUR, CP.
* ST	=	SURFACE TENSION OF LIQUID, dyne/cm.
* A0-A		CONSTANTS FOR TWO-PHASE MODULUS EQUATION
* Hx		HUNTINGTON CORRELATION
* FH		HUNTINGTON FRICTION FACTOR
* XD	=	DAVIS TWO-PHASE CORRELATION=X IN DISPERSED FLOW EQUATION
* P	=	PARACHOR FOR HYDROCARBON
* MW * BX	=	MOLECULAR WEIGHT OF HYDROCARBON
* BX		BAKER'S PARAMETER IN THE LIQUID-PHASE
* BY		BAKER'S PARAMETER IN THE VAPOR-PHASE
* PR		ABSOLUTE ROUGHNESS OF PIPE WALL, ft.
* K		VELOCITY HEAD TO DUE PIPE FITTINGS.
		TWOPHASE FLOW MODULUS.
*****	****	*************************

COMMON/OVID1/ WL, VISL, DENL, ST COMMON/OVID2/ WG, VISG, DENG COMMON/OVID3/ ID, D, PR COMMON/OVID4/ DPST COMMON/OVID5/ DPW COMMON/OVID6/ DPP COMMON/OVID7/ DPSL

COMMON/OVID8/ DPB COMMON/OVID9/ DPAN COMMON/OVID10/ DPD COMMON/OVID11/ DELTPL, DELTPG, DELTP, DELTPT COMMON/OVID12/ X, VEL COMMON/OVID12/ X, VEL COMMON/OVID13/ BX, BY COMMON/OVID14/ REL, LFF, REGG, GFF COMMON/OVID15/ REG COMMON/OVID16/ LEQ, LST, L COMMON AREA REAL ID, L, LEQ, LFF, LST, K c c THE FLOW REGIMES FOR TWO-PHASE FLOWS ARE: Stratified, Wave, Plug, Slug, Bubble, Annular, Dispersed DATA REG /'STRATIFIED', 'WAVE ', 'PL' 'BUBBLE ', 'ANNULAR ', 'DISPERSED '/ 'PLUG '. 'SLUG ٠, * с с OPEN THE DATA FILE. OPEN (UNIT = 3, FILE = 'DATA35.DAT', STATUS = 'OLD', ERR = 18) с OPEN FILE FOR PRINTING THE RESULTS. OPEN (UNIT = 1, FILE = 'PRN') WRITE (1, 100) 100 FORMAT (///,10X,'TWO-PHASE PRESSURE LOSS CALCULATION IN A',\) WRITE (1, 110) FORMAT (' PIPE LINE', /78(1H*)) 110 с READ IN THE DATA IN DATAFILE READ THE PIPE INSIDE DIAMETER, inch: ID READ THE STRAIGHT LENGTH OF PIPE, ft.: LST READ THE LIQUID FLOW RATE, lb/hr.: WL READ THE LIQUID VISCOSITY, cP: VISL READ THE LIQUID DENSITY, lb/ft^3.: DENL READ THE LIQUID SURFACE TENSION, dyne/cm.: ST READ THE GAS FLOW RATE, lb/hr.: WG. READ THE GAS VISCOSITY, cP: VISG. DEAD THE GAS VISCOSITY, cP: VISG. с с с с с с с с с READ THE GAS DENSITY, 1b/ft^3.: READ THE PIPE ROUGHNESS, ft.: PR1 с READ THE VELOCITY HEAD DUE TO PIPE FITTINGS (e.g. bends, valves c c otherwise give the default value of zero): K READ (3, *, ERR=19) ID, LST, WL, VISL READ (3, *, ERR=19) DENL, ST, WG, VISG READ (3, *, ERR=19) DENG, PR1, K

GO TO 20

```
WRITE (*, 120)
FORMAT (5X, 'DATA FILE DOES NOT EXIST')
18
120
       GO TO 1000
19
       WRITE (*, 130)
       FORMAT (5x, 'ERROR MESSAGE IN THE DATA VALUE')
130
       GO TO 1000
20
       LEQ = 0.0
       FACTOR = 0.0
       PR = 0.00015
       PI = 3.1415927
с
       CALCULATE THE PIPE AREA, ft^2.
       D = ID/12.0
       AREA = (PI*D*D)/4.0
с
       CHECK WHETHER THE PIPE ROUGHNESS IS CARBON STEEL (i.e. PR=0.00015 ft)
       IF (PR .NE. PR1) THEN
       PR = PR1
       ENDIF
       DETERMINE THE FLOW REGIME FROM THE BAKER MAP
с
       CALL BAKER
       CALCULATE THE EQUIVALENT LENGTH OF PIPE DUE TO FITTINGS, ft.
С
       LEQ = (K*D)/LFF
с
       CALCULATE THE TOTAL LENGTH OF PIPE, ft.
       L = LST + LEQ
       С
c
c
       * CONSIDER WHETHER PIPE EROSION/CORROSION MAY OCCUR *
с
       CALCULATE THE SUPERFICIAL LIQUID AND GAS VELOCITIES, ft/sec.
       UL = WL/(DENL*AREA*3600.0)
       UG = WG/(DENG*AREA*3600.0)
с
       CALCULATE VELOCITY OF THE MIXTURE
       UM = UL + UG
с
       CALCULATE DENSITY OF THE MIXTURE
       DENM = (WL + WG)/((WL/DENL) +(WG/DENG))
С
       CALCULATE THE INDEX FOR EROSION
       FACTOR = FACTOR + DENM*(UM**2)
```

```
с
```

```
C1 = 17578.98/(BX**0.6548)
        IF (BY .GT. C1) THEN
GO TO 30
        ELSE
        C2 = 5866.07/(BX**0.1901)
        ENDIF
        IF (BY .GT. C2) THEN
GO TO 35
ENDIF
        *****
С
с
        USING THE REGIME FOR STRATIFIED FLOW, DETERMINE THE TWO-PHASE
с
        MODULUS
        CALL STRAT (YG, REGIME, IR)
              GO TO 999
        USING THE REGIME FOR WAVE FLOW, DETERMINE THE TWO-PHASE MODULUS
с
   35
       CALL WAVE (YG, REGIME, IR)
        GO TO 999
       C5 = 2126105.0/(BX**1.0173)
   30
        IF (BY .GT. C5) THEN
GO TO 40
        ELSE
        C6 = 2491.41/(BX**0.2189)
        ENDIF
        IF (BY .GT. C6) THEN
        .Gr. C6
GO TO 41
ENDIF
с
с
        USING THE REGIME FOR PLUG FLOW, DETERMINE THE TWO-PHASE FLOW
        MODULUS.
        CALL PLUG (YG, REGIME, IR)
              GO TO 999
        C4 = 10.7448 - (1.6265 + ALOG(BX)) + (0.2839 + (ALOG(BX)) + 2)
   41
        C4 = EXP(C4)
IF (BY .GT. C4) THEN
        GO TO 45
        ELSÉ
        ENDIF
c
c
        USING THE REGIME FOR SLUG FLOW, DETERMINE THE TWO-PHASE FLOW
        MODULUS.
        CALL SLUG (YG, REGIME, IR)
GO TO 999
   40
       IF (BX .GT. 150.0) THEN
```

GO TO 50 ELSE GO TO 45 ENDIF USING THE REGIME FOR BUBBLE FLOW, DETERMINE THE TWO-PHASE FLOW C C MODULUS. 50 CALL BUBBLE (YG, REGIME, IR) GO TO 999 C3 = 11.3976 - (0.6084 * ALOG(BX)) + (0.0779 * (ALOG(BX)) * 2)45 C3 = EXP(C3)IF (BY .GT. C3) THEN GO TO 55 ELSE ENDIF USING THE REGIME FOR ANNULUS FLOW, DETERMINE THE TWO-PHASE FLOW C C MODULUS. CALL ANNUL (YG, REGIME, IR) GO TO 999 USING THE REGIME FOR DISPERSED FLOW, DETERMINE THE TWO-PHASE FLOW с ċ MODULUS. 55 CALL DISPER (YG, REGIME, IR) CALL OUTPUT (WL, REL, LFF, DELTPL, WG, REGG, GFF, DELTPG, BX, BY, YG, FACTOR, IR, REGIME, DELTP, DELTPT) 999 CLOSE (3, STATUS = 'KEEP') CLOSE (1) 1000 STOP END 00000 THIS PROGRAM USES THE SEMI-EMPIRICAL METHOD OF LOCKHART-MARTINELLI FOR CALCULATING THE PRESSURE DROP OF EITHER THE LIQUID OR THE GAS PHASE FLOW IN A PIPELINE SUBROUTINE BAKER COMMON/OVID1/ WL, VISL, DENL, ST COMMON/OVID2/ WG, VISG, DENG COMMON/OVID13/ ID, D, PR COMMON/OVID11/ DELTPL, DELTPG, DELTPT, DELTPT COMMON/OVID12/ X, VEL COMMON/OVID13/ BX, BY COMMON/OVID14/ REL, LFF, REGG, GFF COMMON/OVID14/ LEQ, LST, L COMMON/OVID16/ LEQ, LST, L REAL ID, L, LEQ, LST, LFF

```
CALCULATE THE LIQUID REYNOLDS NUMBER THROUGH PIPE AND THE
c
c
     FRICTION FACTOR OF THE PIPE.
     REL = (6.31*WL)/(ID*VISL)
      IF (REL .LE. 2100.0) THEN
          GO TO 20
      ELSE
      LFF = 4.0/B1**2
      ENDIE
     GO TO 30
  20
     LFF = 64.0/REL
     CALCULATE LIQUID PRESSURE DROP OF THE PIPE
С
     DELTPL = (0.000366*LFF*(WL**2))/((ID**5)*DENL)
  30
      **********
С
С
      CALCULATE THE GAS REYNOLDS NUMBER THROUGH PIPE AND THE FRICTION
С
      FACTOR OF THE PIPE.
с
      *****
     REGG = (6.31 \times WG) / (ID \times VISG)
     IF ( REGG .LE. 2100.0) THEN
          GO TO 50
      ELSE
      A2 = ((PR/D)/3.7+(6.7/REGG)**0.9)
      B2 = -4.0*ALOG10((PR/(3.7*D))-(5.02/REGG)*ALOG10(A1))
      GFF = 4.0/B2**2
      ENDIF
     GO TO 60
     GFF = 64.0/REGG
  50
С
     CALCULATE THE GAS PRESSURE DROP OF THE PIPE
  60
     DELTPG = (0.000366*GFF*(WG**2))/((ID**5)*DENG)
с
      C
C
      LOCKHART-MARTINELLI TWO-PHASE FLOW MODULUS
      X = SQRT(DELTPL/DELTPG)
с
     CALCULATE THE VELOCITY OF THE TWO-PHASE MIXTURE IN THE PIPE
      VEL = (0.0509/(ID**2))*((WG/DENG)+(WL/DENL))
С
      CALCULATE BAKER'S PARAMETERS, BX AND BY
C
C
     VAL3 = SQRT(DENL*DENG)/(DENL**(2./3.))
      VAL4 = (VISL**(1./3.))/ST
     BX = 531.0*(WL/WG)*VAL3*VAL4
```

```
BY = 2.16*(WG/AREA)/SORT(DENL*DENG)
        RETURN
        END
С
        THIS PROGRAM CALCULATES THE TWO PHASE FLOW MODULUS AND PRESSURE
С
ċ
        DROPS FOR STRATIFIED FLOW REGIME
с
          SUBROUTINE STRAT (YGST, REGIME, IR)
        CHARACTER*10 REGIME(7),REG(7)
        COMMON/OVID1/ WL, VISL, DENL, ST
COMMON/OVID4/ DPST
        COMMON/OVID1/ DELTPL, DELTPG, DELTP, DELTPT
COMMON/OVID12/ X, VEL
COMMON/OVID15/ REG
COMMON/OVID15/ REG
        COMMON AREA
        REAL LEQ, LST, L
        IR = 1
        VAL5 = 15400.0*X
        VAL6 = (WL/AREA)**0.8
        YGST = (VAL5/VAL6) **2
        DPST = DELTPG*YGST
        DELTP = DELTP*L/100.0
        REGIME(IR) = REG(1)
        RETURN
        END
        С
C
C
C
        THIS PROGRAM CALCULATES THE TWO PHASE FLOW MODULUS AND PRESSURE
        DROPS FOR WAVE FLOW REGIME.
        *****
        SUBROUTINE WAVE (YGW, REGIME, IR)
        CHARACTER*10 REGIME(7),REG(7)
COMMON/OVID1/ WL, VISL, DENL, ST
COMMON/OVID2/ WG, VISG, DENG
COMMON/OVID3/ ID, D, PR
COMMON/OVID5/ DPW
COMMON/OVID11/ DELTPL, DELTPG, DELTP, DELTPT
COMMON/OVID15/ REG
COMMON/OVID16/ LEQ, LST, L
COMMON AREA,FH
        REAL ID, LEQ, LST, L
        IR = 2
        HX = 2
HX = (WL/WG)*(VISL/VISG)
VAL7 = (0.2111*ALOG(HX))-3.993
        FH = EXP(VAL7)
        DPW = (0.000366*FH*(WG**2))/((ID**5)*DENG)
```

YGW = 0.0

```
DELTP = DPW
        DELTPT = DELTP*L/100.0
        REGIME(IR) = REG(2)
        RETURN
        END
         ********
C
C
C
        THIS PROGRAM CALCULATES THE TWO PHASE FLOW MODULUS AND PRESSURE
        DROPS FOR PLUG FLOW REGIME.
         -----
с
        SUBROUTINE PLUG (YGP, REGIME, IR)
CHARACTER*10 REGIME(7),REG(7)
        COMMON/OVID1/ WL, VISL, DENL, ST
COMMON/OVID6/ DPP
COMMON/OVID11/ DELTPL, DELTPG, DELTP, DELTPT
COMMON/OVID12/ X, VEL
COMMON/OVID15/ REG
COMMON/OVID16/ LEQ, LST, L
         COMMON AREA
        REAL LEQ, LST, L
         IR = 3
         VAL8 = 27.315*(X)**0.855
         VAL9 = (WL/AREA)**0.17
         YGP = (VAL8/VAL9) * 2
         DPP = DELTPG*YGP
        DELTP = DPP
        DELTPT = DELTP*L/100.0
        REGIME(IR) = REG(3)
        RETURN
        END
0000
         THIS PROGRAM CALCULATES THE TWO PHASE FLOW MODULUS AND PRESSURE
        DROPS FOR SLUG FLOW REGIME.
         SUBROUTINE SLUG (YGSL, REGIME, IR)
         CHARACTER*10 REGIME(7),REG(7)
         COMMON/OVID1/ WL, VISL, DENL, ST
COMMON/OVID7/ DPSL
        COMMON/OVID11/ DELTPL, DELTPG, DELTP, DELTPT
COMMON/OVID12/ X, VEL
COMMON/OVID15/ REG
COMMON/OVID15/ LEQ, LST, L
        COMMON AREA
        REAL LEQ, LST, L
         TR = 4
         VAL10 = 1190.0*(X)**0.815
         VAL11 = (WL/AREA)**0.5
YGSL = (VAL10/VAL11)**2
DPSL = DELTPG*YGSL
```

```
DELTP = DPSL
        DELTPT = DELTP*L/100.0
        REGIME(IR) = REG(4)
        RETURN
        END
с
        THIS PROGRAM CALCULATES THE TWO PHASE FLOW MODULUS AND PRESSURE
        DROPS FOR BUBBLE FLOW REGIME.
         SUBROUTINE BUBBLE (YGB, REGIME, IR)
        CHARACTER*10 REGIME(7),REG(7)
        CHARACTERTIO RECIME(7),REG(7)
COMMON/OVID1/ WL, VISL, DENL, ST
COMMON/OVID8/ DPB
COMMON/OVID11/ DELTPL, DELTPG, DELTP, DELTPT
COMMON/OVID12/ X, VEL
COMMON/OVID15/ REG
COMMON/OVID16/ LEQ, LST, L
COMMON AREA
        REAL LEQ, LST, L
        IR = 5
        VAL12 = 14.2*(X**0.75)
         VAL13 = (WL/AREA)**0.1
        YGB = (VAL12/VAL13)**2
        DPB = DELTPG*YGB
        DELTP = DPB
        DELTPT = DELTP*L/100.0
        REGIME(IR) = REG(5)
        RETURN
        END
         ******************************
        THIS PROGRAM CALCULATES THE TWO PHASE FLOW MODULUS AND PRESSURE
        DROPS FOR ANNULAR FLOW REGIME.
        SUBROUTINE ANNUL (YGAN, REGIME, IR)
        CHARACTER*10 REGIME(7),REG(7)
        COMMON/OVID3/ ID, D, PR
COMMON/OVID9/ DPAN
        COMMON/OVID1/ DELTPL, DELTPG, DELTP, DELTPT
COMMON/OVID12/ X, VEL
COMMON/OVID15/ REG
COMMON/OVID15/ REG
COMMON/OVID16/ LEQ, LST, L
        REAL ID, LEQ, LST, L
        IR = 6
        IF (ID .GT. 12.0) THEN
              GO TO 5
        ELSE
```

```
C1 = 4.8 - 0.3125 \times ID
           C2 = 0.343 - 0.021 * ID
```

c c č

```
ENDIF
        GO TO 15
       D = 10.0
    5
        C1 = 4.8-0.3125*D
C2 = 0.343-0.021*D
        VAL14 = C1*(X**C2)
YGAN = VAL14**2
DPAN = DELTPG*YGAN
   15
        DELTP = DPAN
        DELTPT = DELTP*L/100.0
        REGIME(IR) = REG(6)
        RETURN
        END
0000
         THIS PROGRAM CALCULATES THE TWO PHASE FLOW MODULUS AND PRESSURE
        DROPS FOR DISPERSED FLOW REGIME.
         ******************
                                               *************
        SUBROUTINE DISPER (YGD, REGIME, IR)
        CHARACTER*10 REGIME(7), REG(7)
        COMMON/OVID10/ DPD
COMMON/OVID11/ DELTPL, DELTPG, DELTP, DELTPT
COMMON/OVID12/ X, VEL
COMMON/OVID15/ REG
COMMON/OVID16/ LEQ, LST, L
        REAL LEQ, LST, L
        IR = 7
        A0 = 1.4659
        A1 = 0.4914
        A2 = 0.0489
         A3 = -0.0003487
        XS = -0:000345
VAL15 = A0+A1*ALOG(X)+A2*(ALOG(X))**2+A3*(ALOG(X))**3
YGD = (EXP(VAL15))**2
DPD = DELTPG*YGD
        DELTP = DPD
        DELTPT = DELTP*L/100.0
        REGIME(IR) = REG(7)
        RETURN
        END
С
        č
c
         * THIS PROGRAM PRINTS THE RESULTS OF THE TWO-PHASE FLOW
        SUBROUTINE OUTPUT (WL, REL, LFF, DELTPL, WG, REGG, GFF, DELTPG,
1 BX, BY, YG, FACTOR, IR, REGIME, DELTP, DELTPT)
        CHARACTER*10 REGIME(7), REG(7)
        COMMON/OVID3/ ID, D, PR
COMMON/OVID12/ X, VEL
COMMON/OVID15/ REG
```

	COMMON/OVID16/ LEQ, LST, L COMMON FH
	REAL ID, LEQ, LST, L, LFF
140 1 2 3	<pre>WRITE (1, 140) ID, LEQ, LST, L FORMAT (5X,'PIPE INTERNAL DIAMETER, inch:', T60, F9.3,/, 5X,'EQUIVALENT LENGTH OF PIPE, ft.:', T60, F9.3,/, 5X,'ACTUAL LENGTH OF PIPE, ft.:', T60, F9.3,/, 5X,'TOTAL LENGTH OF PIPE, ft.:', T60, F9.3)</pre>
150 1 2 3	<pre>WRITE (1, 150) WL, REL, LFF, DELTPL FORMAT (5X,'LIQUID FLOWRATE, lb/hr.:', T60, F12.3,/, 5X,'LIQUID REYNOLDS NUMBER:', T60, F12.0,/, 5X,'LIQUID FRICTION FACTOR:', T60, F8.4,/, 5X,'PRESSURE DROP OF LIQUID, psi/100ft.:', T60,F8.4)</pre>
160 1 2 3	WRITE (1, 160) WG, REGG, GFF, DELTPG FORMAT (5X,'GAS FLOW RATE, lb/hr.:', T60, F12.3,/, 5X,'GAS REYNOLDS NUMBER:', T60, F12.0,/, 5X,'GAS FRICTION FACTOR:', T60, F8.4,/, 5X,'PRESSURE DROP OF GAS, psi/100ft.:', T60, F8.4)
170	WRITE (1, 170) REGIME(IR) Format (5x,'flow regime Is:',T60, A)
180 *	WRITE (1, 180) X FORMAT (5X,'LOCKHART-MARTINELLI TWO PHASE FLOW MODULUS:', T60,F8.4)
190 1 2	WRITE (1, 190) VEL, BX, BY FORMAT (5X,'VELOCITY OF FLUID IN PIPE, ft./sec.:', T60, F8.3,/, 5X,'BAKER PARAMETER IN THE LIQUID PHASE:', T60, F12.3,/, 5X,'BAKER PARAMETER IN THE GAS PHASE:', T60, F12.3)
	IF (YG .EQ. 0.0) THEN
200	WRITE (1, 200) FH FORMAT (5X,'HUNTINGTON FRICTION FACTOR:', T60, F8.4)
	ELSE
210	WRITE (1, 210) YG FORMAT (5X,'TWO-PHASE FLOW MODULUS:', T60, F8.4) ENDIF
220 1 2 3	<pre>WRITE (1, 220) DELTP, DELTPT FORMAT (5X,'PRESSURE DROP OF TWO-PHASE MIXTURE, psi/100ft:', T60, F8.4,/, 5X,'OVERALL PRESSURE DROP OF THE TWO-PHASE MIXTURE, psi:', T60, F8.4)</pre>
	IF (FACTOR .GT. 10000.0) THEN
230	WRITE (1, 230) FACTOR FORMAT (5X,'INDEX', 1X, F10.0, ' IS GREATER THAN 10000',\)
240	WRITE (1, 240) FORMAT (' PIPE EROSION IS POSSIBLE')

	ELSE
250	WRITE (1, 250) FACTOR FORMAT (5%,'INDEX',1%, F10.0, ' IS LESS THAN 10000'\)
260	WRITE (1, 260) FORMAT (' PIPE EROSION IS UNLIKELY') ENDIF
270	WRITE (1, 270) Format (78(1H-))
с	FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR(12)
	RETURN END

PROGRAM PROG36

c c	**************************************		
с с с	TWO-PHASE VERTICAL PIPE. IT FURTHER PRINTS OUT A MESSAGE IF THE PIPE IS SELF-VENTING, IF PULSATION FLOW OCCURS, OR IF NO PRESSURE GRADIENT IS NEEDED.		
0000	AREA = INSIDE CROSS-SECTIONAL AREA OF PIPE, ft^2 .		
c c	FRNL = FROUDE NUMBER FOR LIQUID PHASE FRNG = FROUDE NUMBER FOR GAS PHASE		
C C C	G = GRAVITATIONAL CONSTANT (32.2 ft/s^2) DENL = LIQUID DENSITY, lb/ft^3 VL = LIQUID VELOCITY, ft/s. VG = VAPOR VELOCITY, ft/s.		
C C C	WL = LIQUID FLOWRATE, ID/h. WG = VAPOR FLOWRATE, lb/h.		
с	************************		
С	OPEN FILES TO READ THE DATA AND TO PRINT THE RESULTS		
	PARAMETER (G = 32.2, PI = 3.1415927)		
с	OPEN THE DATA FILE.		
	OPEN (UNIT = 3, FILE = 'DATA36.DAT', STATUS = 'OLD', ERR = 18)		
с	OPEN FILE FOR PRINTING THE RESULTS.		
	OPEN (UNIT = 1, FILE = 'PRN')		
100 *	<pre>WRITE (1, 100) FORMAT (///,15X, 'VAPOR-LIQUID TWO-PHASE VERTICAL DOWNFLOW',/1H ,</pre>		
с	READ THE INPUT DATA		
00000	READ THE PIPE INSIDE DIAMETER, inch: D READ THE LIQUID FLOW RATE, lb/hr.: WL READ THE LIQUID DENSITY, lb/ft^3: DENL READ THE GAS FLOW RATE, lb/hr.: WG. READ THE GAS DENSITY, lb/ft^3.: DENG.		
	READ (3, *, ERR=19) D, WL, DENL READ (3, *, ERR=19) WG, DENG GO TO 10		
18 110	WRITE (*, 110) Format (5x, 'Data file does not exist')		
	GO TO 999		

19 120	WRITE (*, 120) FORMAT (5X,'ERROR MESSAGE IN THE DATA VALUE') GO TO 999
10	D1 = D/12. AREA = (PI*D1**2)/4.0
130	WRITE (1, 130) D FORMAT (10X, 'PIPE INTERNAL DIAMETER, inch:', T65, F8.3)
140 *	WRITE (1, 140) WL, DENL FORMAT (10X, 'LIQUID FLOWRATE, lb/h.:', T65, F8.0, /, 10X, 'LIQUID DENSITY, lb/ft^3:', T65, F8.3)
150 *	WRITE (1, 150) WG, DENG FORMAT (10X, 'VAPOR FLOWRATE, lb/h.:', T65, F8.0, /, 10X, 'VAPOR DENSITY, lb/ft^3:', T65, F8.3)
160	WRITE (1, 160) AREA FORMAT (10X, 'PIPE AREA, ft^2:', T65, F8.3)
с	CALCULATE LIQUID AND VAPOR VELOCITIES
	VL = WL/(3600.0*DENL*AREA) VG = WG/(3600.0*DENG*AREA)
170 *	WRITE (1, 170) VL, VG FORMAT (10X, 'LIQUID VELOCITY, ft/s:', T65, F8.3, /, 10X, 'VAPOR VELOCITY, ft/s:', T65, F8.3)
с	CALCULATE THE LIQUID PHASE AND THE VAPOR PHASE FROUDE NUMBERS
	<pre>FRNL = (VL/((G*D1)**0.5))*(DENL/(DENL-DENG))**0.5 FRNG = (VG/((G*D1)**0.5))*(DENG/(DENL-DENG))**0.5</pre>
180 *	WRITE (1, 180) FRNL, FRNG FORMAT (10X, 'FROUDE NUMBER FOR LIQUID PHASE:', T65, F8.4, /, 10X, 'FROUDE NUMBER FOR VAPOR PHASE:', T65, F8.4)
с	TEST FOR SELF-VENTING, PULSE FLOW OR NO PRESSURE GRADIENT.
	IF (FRNL .LT. 0.31) THEN
190	WRITE (1, 190) FORMAT (10X, 'LINE IS SELF-VENTING THEREFORE NO VIBRATION',\)
200	WRITE (1, 200) FORMAT (' PROBLEMS ARE EXPECTED') ELSEIF (FRNL .GE. 0.31 .AND. FRNL .LT. 1.0) THEN
210	WRITE (1, 210) FORMAT (10X, 'FLOW IS PULSE AND THIS MAY RESULT IN PIPE')
220	WRITE (1, 220) FORMAT (' VIBRATION') ELSEIF (FRNL .GT. 1.0) THEN
	WRITE (1, 230)

230	FORMAT (10X, 'NO PRESSURE GRADIENT') ENDIF	
240	WRITE (1, 240) Format (78(1H-))	
с	FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.	
	WRITE (1, *) CHAR(12)	
	CLOSE (3, STATUS = 'KEEP') CLOSE (1)	
999	STOP END	

PROGRAM PROG37

	 THIS PROGRAM CALCULATES LINE SIZES FOR FLASHING STEAM CONDENSATE. THE PROGRAM WILL COMPUTE THE FOLLOWING: 1. CALCULATES THE AMOUNT OF CONDENSATE FLASHED FOR ANY GIVEN CONDENSATE HEADER PRESSURE FROM 15 TO 140 psia. AND INITIAL STEAM PRESSURES MAY VARY BETWEEN 40 AND 165 psia. 2. THE RETURN CONDENSATE HEADER TEMPERATURE. 3. THE PRESSURE DROP (psi/100ft.) OF THE STEAM CONDENSATE MIXTURE IN THE RETURN HEADER. 4. THE VELOCITY OF THE STEAM CONDENSATE MIXTURE AND 5. INDICATES A WARNING MESSAGE IF VELOCITY IS EQUAL TO OR GREATER THAN 5000 ft/min. 		
c c	NOMENCLATURE.		
000000000000000000000000000000000000000	<pre>D = PIPE INTERNAL DIAMETER, inch. DELTP = PRESSURE DROP OF FLASHED CONDENSATE MIXTURE,</pre>		
100 *	<pre>WRITE (1, 100) FORMAT (///, 15X, 'LINE SIZING FOR FLASHING STEAM CONDENSATE',/, 78(1H*))</pre>		
с	READ THE INPUT DATA		
с с с с	READ THE PIPE INTERNAL DIAMETER, inch: D READ THE FLUID FLOW RATE, lb/hr.: W READ THE CONDENSATE STEAM PRESSURE BEFORE FLASHING, psia.: PC READ THE FLASHED CONDENSATE HEADER PRESSURE, psia.: PH		

READ (3, *, ERR=19) D, W READ (3, *, ERR=19) PC, PH GO TO 10 WRITE (*, 110) FORMAT (10X, 'DATA FILE DOES NOT EXIST') 18 110 GO TO 999 19 WRITE (*, 120) FORMAT (10X, 'ERROR MESSAGE IN THE DATA VALUE') 120 GO TO 999 10 X = 6.122 - (16.919/ALOG(PH))A = 0.00671*(ALOG(PH))**2.27 B = EXP(X) * (10 * * (-4)) + 0.0088CALCULATE WEIGHT FRACTION OF CONDENSATE FLASHED TO VAPOR. С WFRFL = B*(ALOG(PC))**2 - ACALCULATE FLASHED STEAM FLOWRATE AND FLASHED STEAM С С CONDENSATE LIQUID FLOWRATE, 1b/h. WG = WFRFL*W WL = W - WGWRITE (1, 130) D, W FORMAT (10X, 'PIPE INTERNAL DIAMETER, inch:', T65, F8.3, /, 10X,'TOTAL FLOW OF MIXTURE IN CONDENSATE HEADER, 1b/h:', 130 * T65, F8.0) WRITE (1, 140) PC, PH FORMAT (10X,'STEAM CONDENSATE PRESSURE BEFORE FLASHING, psia:', 765, F8.3, /, 10X,'FLASHED CONDENSATE HEADER PRESSURE, psia:', 140 T65, F8.3) WRITE (1, 150) WFRFL FORMAT (10X,'WEIGHT FRACTION OF CONDENSATE FLASHED TO VAPOR:', * T65, F8.3) 150 WRITE (1, 160) WG, WL FORMAT (10X, 'FLASHED STEAM FLOWRATE, 1b/h.:', T65, F8.0,/, * 10X,'FLASHED CONDENSATE LIQUID FLOWRATE, 1b/h.:', T65, F8.0) 160 CALCULATE THE TEMPERATURE OF FLASHED CONDENSATE. С TFL = 115.68*(PH)**0.226CALCULATE THE FLASHED STEAM DENSITY, FLASHED CONDENSATE LIQUID ¢ c DENSITY AND DENSITY OF MIXTURE (FLASHED CONDENSATE/STEAM) 1b/ft^3 DENG = 0.0029*(PH)**0.938 DENL = 60.827 - (0.078*PH) + (0.00048*(PH)**2) - (0.0000013*(PH)**3)

DENMIX = W/(WG/DENG + WL/DENL)

FOR TURBULENT FLOW, CALCULATE THE FRICTION FACTOR. с FF = 0.25/(-ALOG10(0.000486/D))**2DELTP = (0.000336*FF*W**2)/(DENMIX*D**5)CALCULATE THE VELOCITY OF FLASHED CONDENSATE MIXTURE, ft/min. C VEL = (3.054/D**2)*(WG/DENG + WL/DENL)WRITE (1, 170) TFL, DENG FORMAT (10X, 'TEMPERATURE OF FLASHED CONDENSATE, oF:', T65, F8.3, /, 10X, 'FLASHED STEAM DENSITY, 1b/ft^3:', T65, F8.3) 170 WRITE (1, 180) DENL, DENMIX FORMAT (10X, 'FLASHED CONDENSATE LIQUID DENSITY, 1b/ft^3:', * T65, F8.3, /, 10X, 'DENSITY OF MIXTURE, 1b/ft^3:', T65, F8.3) 180 WRITE (1, 190) FF, DELTP, VEL FORMAT (10X, 'FRICTION FACTOR:', T65, F8.4, /, 10X, 'PRESSURE DROP OF FLASHED, psi:',/,10X, 'CONDENSATE MIXTURE, psi/100ft:', T65, F8.3,/,10X, 'VELOCITY OF FLASHED CONDENSATE MIXTURE, ft/min.:', %CF, T0000 190 * * ٠ T65, F10.0) DETERMINE WHETHER PROCESS LINE MAY DETERIORATE i.e. WHETHER с č VELOCITY IS EQUAL OR GREATER THAN 5000 ft./min. IF (VEL .GE. 5000.0) THEN WRITE (1, 200) 200 FORMAT (10X, 'FLASHED CONDENSATE MIXTURE VELOCITY IS GREATER', \) WRITE (1, 210) FORMAT (' THAN 5000 ft./min.') 210 WRITE (1, 220) FORMAT (10X, 'DETERIORATION OF THE LINE IS POSSIBLE') 220 ELSE WRITE (1, 230) 230 FORMAT (10X, 'FLASHED CONDENSATE MIXTURE IS LESS THAN', \) WRITE (1, 240) FORMAT (' 5000 ft/min.') 240 WRITE (1, 250) FORMAT (10X, 'THE CONDENSATE HEADER LINE WILL NOT DETERIORATE') 250 ENDIF WRITE (1, 260) FORMAT (78(1H-)) 260 С FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE. WRITE (1, *) CHAR(12) CLOSE (3, STATUS = 'KEEP') CLOSE (1) 999 STOP END

с с с с с с	THIS PROGRAM CALCULATES THE TOTAL PRESSURE DROP IN A PACKED BED WHEN LIQUID OR VAPOR STREAMS ARE PASSED THROUGH IT. THE PROGRAM ALSO CALCULATES THE FLUID REYNOLDS NUMBER AND THE FRICTION FACTOR.		
с с с с с с с	NOMENCLATURE A = CROSS-SECTIONAL AREA OF BED, ft^2. AP = AREA OF PARTICLES, ft^2. BD = PACKED BED DIAMTER, ft.		
с с с с с	BL = PACKED BED LENGTH, ft. DEN = DENSITY OF FLUID AT FLOWING CONDITIONS, lb/ft^3. DELPT = TOTAL PRESSURE DROP IN PACKED BED, lb/in^2. FV = FRACTION VOIDS IN PACKED BED. FP = FRICTION FACTOR.		
0000	GC = GRAVITATIONAL CONSTANT = 4.17 x 10^8 (lbm/lbf)(ft/h^2) MW = MOLECULAR WEIGHT OF FLUID. P = FLUID PRESSURE, psia. PD = PARTICLE DIAMETER, inch. PL = PARTICLE INGTH inch.		
0 0 0 0 0 0 0 0	RE = REYNOLDS NUMBER S = PACKED BED SURFACE AREA, (ft^2/ft^3 BED). T = FLUID TEMPERATURE, oR. VIS = FLUID VISCOSITY, cP. W = FLUID FLOWRATE, lb/h. Z = COMPRESSIBILITY FACTOR (Z=1.0)		
	PARAMETER (PI = 3.1415927, GC = 4.17*10**8)		
С	OPEN FILES TO READ THE DATA AND TO PRINT THE RESULTS. OPEN (UNIT = 3, FILE ='DATA38.DAT', STATUS = 'OLD', ERR = 18) OPEN (UNIT = 1, FILE = 'PRN')		
100 *	WRITE (1, 100) FORMAT (///, 25x,'PRESSURE DROP IN A PACKED BED',/1H , 78(1H*))		
с	READ THE DATA VALUES.		
0000000	READ THE PACKED BED DIAMERTER, ft.: BD READ THE PACKED BED LENGTH, ft.: BL READ THE FLUID FLOW RATE, lb/hr.: W READ THE PARTICLE DIAMETER, inch: PD READ THE PARTICLE LENGTH, inch: PL READ THE PARTICLE LENGTH, inch: PL READ THE FRACTION OF VOIDS IN PACKED BED, inch: FV READ THE DENSITY OF FLUID AT FLOWING CONDITION, lb/ft^3.: DEN		

с READ THE FLUID VISCOSITY, CP: VIS READ (3, *, ERR=19) BD, BL, W READ (3, *, ERR=19) PD, PL, FV READ (3, *, ERR=19) DEN, VIS GO TO 10 WRITE (*, 110) FORMAT (10X, 'DATA FILE DOES NOT EXIST') 18 110 GO TO 999 WRITE (*, 120) FORMAT (10%, 'ERROR MESSAGE IN THE DATA VALUE') 19 120 GO TO 999 10 130 140 150 с CALCULATE THE CROSS-SECTIONAL AREA OF BED, ft^2. A = (PI*BD**2)/4.0С CALCULATE THE SUPERFICIAL MASS VELOCITY, W/A G = W/Aс CALCULATE THE AREA OF PARTICLES AND THE VOLUME OF PARTICLES. AP = (1.0/144.0)*((PI*PD**2)/2.0 + (PI*PD*PL))VP = (1.0/1728.0)*(((PD**2)*PI*PL)/4.0) с CALCULATE THE PACKED BED SURFACE AREA, ft^2/ft^3. BED. S = AP * (1 - FV) / VPCALCULATE THE EFFECTIVE PARTICLE DIAMETER, ft. С DP = 6.0*(1-FV)/SCALCULATE THE FLUID REYNOLDS NUMBER, RE с

RE = (G*DP)/(2.419*VIS*(1.0 - FV))

160 * *	<pre>WRITE (1, 160) A, G FORMAT (10X, 'CROSS-SECTIONAL AREA OF BED, ft^2:', T60, F8.4,</pre>
170 *	WRITE (1, 170) AP, VP FORMAT (10X, 'AREA OF PARTICLES, ft^2:', T60, E12.4, /, 10X, 'VOLUME OF PARTICLES, ft^3:', T60, E12.4)
180 *	WRITE (1, 180) S, DP FORMAT (10X, 'THE PACKED BED SURFACE AREA, ft^2/ft^3 BED:', T60, F8.3, /, 10X,'EFFECTIVE PARTICLE DIAMETER, ft.:',T60, F8.4)
190	WRITE (1, 190) RE FORMAT (10X, 'FLUID REYNOLDS NUMBER:', T60, F10.0)
c c	DETERMINE THE FRICTION FACTOR FOR LAMINAR, INTERMEDIATE OR TURBULENT FLOW.
	IF (RE .LT. 1.0) THEN
	FP = 150.0/RE
	WRITE (1, 200) FP
200	FORMAT (10X, 'FRICTION FACTOR FOR LAMINAR FLOW:', T60, F8.4) ELSEIF (RE .GT. 1.0 .AND. RE .LT. 10**4) THEN
	FP = 150.0/RE+1.75
210 *	WRITE (1, 210) FP FORMAT (10X,'FRICTION FACTOR FOR INTERMEDIATE FLOW:', T60, F8.4)
	ELSEIF (RE .GT. 10**4) THEN FP = 1.75
220	WRITE (1, 220) FORMAT (10X,'FRICTION FACTOR FOR TURBULENT FLOW:', T60, F8.2)
	ENDIF
с	CALCULATE THE TOTAL PRESSURE DROP IN PACKED BED, DELTP.
	VAL1 = (BL*(1-FV)*G**2)/(144.0*(FV**3)*DP*GC*DEN) VAL2 = ((150.0*2.419*VIS*(1.0-FV)/(DP*G))+1.75) DELTP = VAL1 * VAL2
230 *	WRITE (1, 230) DELTP FORMAT (10X, 'TOTAL PRESSURE DROP IN PACKED BED, lb/in^2:', T60, F8.3)
240	WRITE (1, 240) FORMAT (78(1H-))
с	FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR(12)
	CLOSE (3, STATUS = 'KEEP') CLOSE (1)
999	STOP END

CHAPTER 4

Equipment Sizing

INTRODUCTION

The process engineer is often required to design separators or knockout drums for removing liquids from process gas streams. This chapter reviews the design of horizontal and vertical separators and the sizing of partly filled horizontal and cylindrical vessels. The chapter concludes by reviewing the sizing of a cyclone and a solid-desiccant gas dryer.

Vessels used for processing in the chemical process industries (CPI) are principally of two kinds: those that are without internals and those with internals. Empty separators are drums that provide intermediate storage or surge of a process stream for a limited or extended period. Alternatively, they provide phase separation by settling.

The second category comprises equipment such as reactors, mixers, distillation columns, and heat exchangers. In some cases, it is important to separate liquid and gas flowing simultaneously through a pipe. This is because the conditions of the flowing mixture and the efficiency of separation may vary widely. Therefore, a separator for such duty must be adequate. In addition, there are constraints due to space or weight that often affect the choice of separators, the need to handle solids or effect a three-phase separation, and the requirements for liquid hold-up. In practice, most separation problems are solved by the following types of separating equipment.

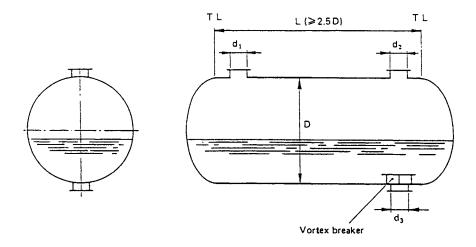
Knockout or Surge Drums (Figures 4-1 and 4-2). A knockout drum is suitable for a bulk separation of gas and liquid, particularly when the liquid volume fraction is high with stratified or plug flow in the pipe. Also, it is useful when vessel internals are required to be kept to a

minimum, e.g., in relief systems or in fouling service. It is unsuitable if a mist is being separated or if high separating efficiency is required.

Cyclones (Figure 4-3). These are robust and not susceptible to fouling or wax. Multicyclones are compact and are reasonably efficient for foam, but not for slugs. The efficiency of a cyclone decreases with increasing diameter, and cyclones are not applicable above 5 ft. diameter. They possess a good efficiency for smaller flow rates. However, cyclones are expensive to operate, especially under vacuum conditions, because they have a higher pressure drop than either a knockout drum or a demister separator.

Demister Separators. A demister separator is fitted either with a vane demister package or with a wire mesh demister mat. The latter type is much preferred, although it is unsuitable for fouling service. The wire mesh demister is a widely applied type of separator, and is adequate for all gas-liquid flow regimes over a wide range of gas flow rates.

A knockout drum or demister separator may be either a vertical or a horizontal vessel. A vertical vessel is generally preferred because its efficiency does not vary with the liquid level. Alternatively, a horizontal



Note: TL = Tangent line



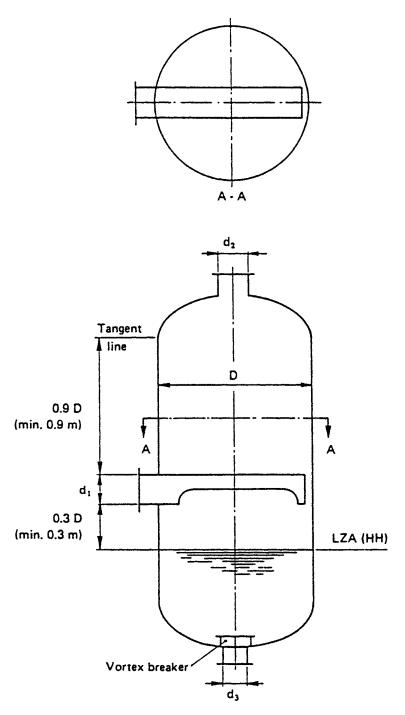


Figure 4-2. Vertical knock-out drum.

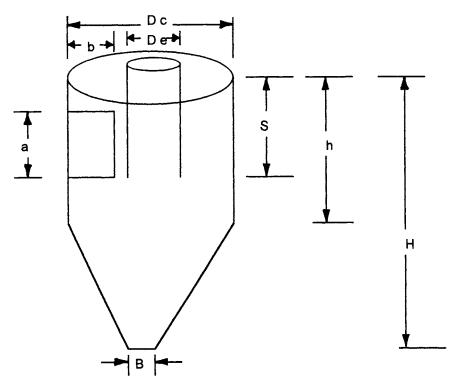


Figure 4-3. A cyclone: design configurations.

vessel is chosen when it offers a clear size advantage, if the headroom is restricted or if a three-phase separation is required. Knockout drums and cyclones are recommended for waxy and coking feeds. Demister mats are not suitable because of the danger of plugging. Vane demister packages are used as alternatives, but provision should be made for cleaning.

SIZING OF VERTICAL AND HORIZONTAL SEPARATORS

Vertical Separators

Vertical liquid-vapor separators are used to disengage a liquid from a vapor when the volume of liquid is small compared with that of vapor volume. The maximum allowable vapor velocity in a vertical separator that reduces the liquid carryover is dependent on the following:

- liquid and vapor densities.
- a constant K based on surface tension, droplet size, and physical characteristics of the system.

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The proportionality constant, K, is 0.35 for oil and gas systems with at least 10-inch disengaging height between the mist eliminator bottom and gas liquid interface. For vertical vessels, K can vary between 0.1 and 0.35, if mist eliminators (demisters) are used to enhance disentrainment. The value of K also depends on the operating pressure of the vessel. At pressures above 30 psig, the K value decreases with pressure with an approximate value of 0.30 at 250 psig and 0.275 at 800 psig. Watkins [1] has developed a correlation between the separation factor and K. Figure 4-4 illustrates Watkin's vapor velocity factor chart, based on five percent of the liquid being entrained with the vapor.

Blackwell [2] has developed a polynomial equation using Watkin's data to calculate the K value for a range of separation factors between 0.006 and 5.0. Watkins proposed a method for sizing reflux drums based upon several factors, as illustrated in Tables 4-1 and 4-2. Table 4-1 gives the recommended design surge times. Table 4-2 gives the multiplying factors for various operator efficiencies. The operating factor is based upon the external unit and its operation, its instrumentation and response to control, the efficiency of labor, and chronic mechanic problems, and the possibility of short or long term interruptions [1].

The multiplying factors F_1 and F_2 represent the instrument and the labor factors.

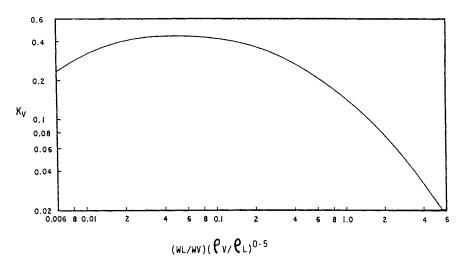


Figure 4-4. Design vapor velocity factor for vertical vapor-liquid separators at 85% of flooding.

			Minutes		
	Instrumen	t Factor F ₁	Labor F	actor F ₂	
Operation	w/Alarm	w/o Alarm	Good	Fair	Poor
FRC	$\frac{1}{2}$	1	1	1.5	2
LRC	1	$1 \frac{1}{2}$	1	1.5	2
TRC	$1\frac{1}{2}$	2	1	1.5	2

 Table 4-1

 Design Criteria for Reflux Distillate Accumulators

 Table 4-2

 Operation Factors for External Units

Operating Characteristics	Factor F ₃
Under good control	2
Under fair control	3
Under poor control	4
Feed to or from storage	1.25
	Factor F ₄
Board mounted level recorder	1.0
Level indicator on board	1.5
Gauge glass at equipment only	2.0

A multiplying factor, F_3 , is applied to the net overhead product going downstream. F_4 depends on the kind and location of level indicators.

It is recommended that 36 inches plus one-half the feed nozzle, OD, (48 inches minimum) be left above the feed nozzle for vapor. Below the feed nozzle, allowance of 12 inches plus one-half the feed nozzle, OD, is required for clearance between the maximum liquid level and the feed nozzle (minimum of 18 inches). At some value between L/D ratios of 3 and 5, a minimum vessel weight will occur resulting in minimum costs for the separator. Figure 4-5 shows the dimensions of a vertical separator.

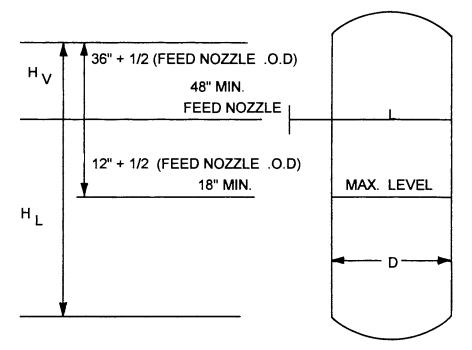


Figure 4-5. Dimension of a vertical separator.

Calculation Method for a Vertical Drum

The following steps are used to size a vertical drum.

Step 1. Calculate the vapor-liquid separation factor.

S. Fac =
$$\frac{WL}{WV} \left(\frac{\rho_v}{\rho_L}\right)^{0.5}$$
 (4-1)

Step 2. From Blackwell's correlation, determine the design vapor velocity factor, K_v , and the maximum design vapor velocity.

$$X = \ln(S. Fac) \tag{4-2}$$

$$K_{v} = \exp(B + DX + EX^{2} + FX^{3} + GX^{4})$$
(4-3)

where B = -1.877478 D = -0.814580 E = -0.187074 F = -0.014523G = -0.001015

$$V_{max} = K_{v} \left(\frac{\rho_{L} - \rho_{v}}{\rho_{v}}\right)^{0.5}$$
(4-4)

Step 3. Calculate the minimum vessel cross-sectional area.

$$Q_v = \frac{WW}{3600.\,\rho_v}, \quad \text{ft}^3/\text{s}$$
 (4-5)

$$A_{v} = \frac{Q_{v}}{V_{max}}, \quad ft^{2}$$
(4-6)

Step 4. Set a vessel diameter based on 6-inch increments and calculate cross-sectional area.

$$D_{\min} = \left(\frac{4.A_{\nu}}{\pi}\right)^{0.5}, \quad \text{ft}$$
(4-7)

 $D = D_{min}$ to next largest 6 inch

Area =
$$\frac{\pi . D_{\min}^2}{4}$$
 (4-8)

Step 5. Estimate the vapor-liquid inlet nozzle based on the following velocity criteria.

$$(U_{max})_{nozzle} = \frac{100}{\rho_{mix}^{0.5}}$$
 ft/s (4-9)

$$(U_{\min})_{\text{nozzle}} = \frac{60}{\rho_{\min}^{0.5}} \quad \text{ft/s}$$
 (4-10)

where

$$\rho_{mix} = \left(\frac{WL + WV}{\frac{WL}{\rho_L} + \frac{WV}{\rho_V}}\right) \frac{lb_m}{ft^3}$$
(4-11)

Step 6. From Figure 4-5, make a preliminary vessel sizing for the height above the center line of a feed nozzle to top seam. Use 36 inches + 1/2 feed nozzle, OD, or 48 inches minimum. Use 12 inches + 1/2 feed

nozzle, OD, or 18 inch minimum to determine the distance below the center line of the feed nozzle to the maximum liquid level.

Step 7. From Table 4-1 or 4-2, select the appropriate full surge volume in seconds. Calculate the required vessel volume.

$$Q_{L} = \frac{WL}{3600.\,\rho_{L}} \qquad ft^{3}/s \tag{4-12}$$

$$V = (Q_L) (\text{design time to fill}), \quad \text{ft}^3$$

= (60.0)(Q₁)(T), \quad ft^3 (4-13)

The liquid height is:

$$H_{L} = V.\left(\frac{4}{\pi . D^{2}}\right), \quad ft$$
(4-14)

Step 8. Check geometry. $(H_L + H_v)/D$ must be between 3 and 5. For small volumes of liquid, it may be necessary to provide more liquid surge than is necessary to satisfy the L/D > 3. If the required liquid surge volume is greater than that obtained in a vessel having L/D < 5, a horizontal drum must be provided.

Calculation Method for a Horizontal Drum

Horizontal vessels are used for substantial vapor-liquid separation where the liquid holdup space must be large. Maximum vapor velocity and minimum vapor space are determined as in the vertical drum, except that $K_{\rm H}$ for horizontal separators is generally set at 1.25K_v. The following steps are carried out in sizing horizontal separators.

Step 1. Calculate the vapor-liquid separation factor by Equation 4-1, and K_v by Equations 4-2 and 4-3.

Step 2. For horizontal vessels

$$K_{\rm H} = 1.25 K_{\rm v}$$
 (4-15)

Step 3. Calculate the maximum design vapor velocity.

$$(U_v)_{max} = K_H \left(\frac{\rho_L - \rho_V}{\rho_V}\right)^{0.5} \quad \text{ft/s}$$
(4-16)

Step 4. Calculate the required vapor flow area.

$$(A_v)_{min} = \frac{Q_v}{(U_v)_{max}} ft^2$$
 (4-17)

Step 5. Select appropriate design surge time from Table 4-1 or Table 4-2, and calculate full liquid volume by Equations 4-12 and 4-13.

The remainder of the sizing procedure is carried out by trial and error as follows:

Step 6. When the vessel is full, the separator vapor area can be assumed to occupy only 15 to 25 percent of the total cross-sectional area. Here, a value of 20 percent is used and the total cross-sectional area is expressed as:

$$(A_{total})_{min} = \frac{(A_v)_{min}}{0.2}$$
 (4-18)

$$D_{\min} = \left(\frac{4.(A_{\text{total}})_{\min}}{\pi}\right)^{0.5}$$
(4-19)

Step 7. Assume the length-to-diameter ratio of 3 (i.e, L/D = 3). Calculate the vessel length.

$$L = (3)(D_{\min})$$
(4-20)

Step 8. Because the vapor area is assumed to occupy 20 percent of the total cross-sectional area, the liquid area will occupy 80 percent of the total cross-sectional area.

$$A_{L} = (0.8)(A_{total})_{min}, \quad ft^{2}$$
 (4-21)

Step 9. Calculate the vessel volume.

$$V_{\text{VES}} = (A_{\text{total}})_{\min}(L), \quad \text{ft}^3$$
(4-22)

Step 10. Calculate liquid surge time.

$$T = \frac{(60.0)(A_{L})(L)(\rho_{L})}{WL} min.$$
(4-23)

Liquid Holdup and Vapor Space Disengagement

The dimensions of both vertical and horizontal separators are based on rules designed to provide adequate liquid holdup and vapor disengaging space. For instance, the desired vapor space in a vertical separator is at least $1\frac{1}{2}$ times the diameter, with 6 inches as the minimum above the top of the inlet nozzle. In addition, a 6-inch minimum is required between the maximum liquid level and the bottom of the inlet nozzle. For a horizontal separator, the minimum vapor space is equal to 20 percent of the diameter or 12 inches.

Wire Mesh Pad

Pads of fine wire mesh induce coalescence of impinging droplets into larger ones, which then separate freely from the gas phase. No standard equations have been developed for the pressure drop across wire mesh because there are no standardized mesh pads. However, as a rule of thumb, the pressure drop of a wire mesh is $\Delta P = 1.0$ in H₂O. Every manufacturer makes a standard high efficiency, very-high efficiency, or high-throughput mesh under various trade names, each for a specific requirement.

Standards for Horizontal Separators

The following specifications are generally standard in the design of horizontal separators [3]:

- 1. The maximum liquid level shall provide a minimum vapor space height of 15 inches, but should not be below the center line of the separator.
- 2. The volume of dished heads is not considered in vessel sizing calculations.
- 3. The inlet and outlet nozzles shall be located as closely as practical to the vessel tangent lines. Liquid outlets shall have anti-vortex baffles.

Piping Requirements

Pipes that are connected to and from the process vessels must not interfere with the good working of the vessels. Therefore, the following guidelines should be observed:

• There should be no valves, pipe expansions or contractions within 10-pipe diameters of the inlet nozzle.

• There should be no bends within 10-pipe diameters of the inlet nozzle except:

for knockout drums and demisters, a bend in the feed pipe is permitted if this is in a vertical plane through the axis of the feed nozzle.

for cyclones, a bend in the feed pipe is allowed if this is in a horizontal plane and the curvature is in the same direction as the cyclone vortex.

- A pipe reducer may be used in the vapor line leading from the separator, but it should be no nearer to the top of the vessel than twice the outlet pipe diameter.
- A gate or ball type valve that is fully opened in normal operation should be used, where a valve in the feed line near the separator cannot be avoided.
- High pressure drops that cause flashing and atomization should be avoided in the feed pipe.
- If a pressure reducing valve in the feed pipe cannot be avoided, it should be located as far upstream of the vessel as practicable.

Nomenclature

A = cross-sectional area of vertical separator, ft^2

- A_L = cross-sectional area of horizontal vessel occupied by liquid phase, ft²
- A_T = total cross-sectional area of horizontal vessel, ft²
- $A_v =$ minimum cross-sectional area of vertical separator, ft²
- D = vessel diameter, ft
- H_L = liquid height, ft
- $H_v = vapor height, ft$
- K = proportionality constant
- $K_{\rm H}$ = vapor velocity factor for horizontal separator
- $K_v =$ vapor velocity factor for vertical separator

L = horizontal vessel level, ft

- S.Fac = separation factor
 - $Q_{\rm L}$ = liquid volumetric flow, ft³/s
 - $Q_v =$ vapor volumetric flow, ft³/s
 - T =liquid surge time, min.
 - V_{max} = maximum vapor velocity, ft/s
 - V_{VES} = horizontal vessel volume, ft³
 - WL = liquid flow rate, lb/h
 - WV = vapor flow rate, lb/h

X = ln (S.Fac) $\rho_L = liquid density, lb/ft^3$ $\rho_v = vapor density, lb/ft^3$

SIZING OF PARTLY FILLED VESSELS AND TANKS

Cylindrical vessels and horizontal tanks are used for the storage of fluids in the chemical process industries. Various level instruments are employed to determine the liquid level in process vessels. The exact liquid volume can be obtained either by calibration of the vessels or by tedious calculations. Partial volumes for horizontal, cylinders with flat, dished, elliptical, and hemispherical ends, and for vertical cylinders are employed for storing process fluids.

Kowal [4] presented charts that allow designers to determine the volume of vessels for different dimensions. In addition, parameters for tray designs and tray layout in distillation columns, such as chord length and circle area, are computed in this chapter.

The Equations

Calculate the area of the segment of a circle from Figure 4-6:

$$\cos(\alpha) = \frac{R - H}{R}$$
(4-24)

$$\alpha = \cos^{-1}\left(\frac{R-H}{R}\right)$$
(4-25)

Area of the triangle:

Area of
$$\Delta = (R - H)(2RH - H^2)$$
(4-26)

Therefore, area of the segment of a circle from Figure 4-6 is:

$$A_{s} = R^{2} Cos^{-1} \left(\frac{R - H}{R} \right) - (R - H)(2RH - H^{2})^{0.5}$$
(4-27)

Figure 4-7 is a nomograph based upon the area of the segment of a circle [5]. The nomograph converts the height (rise) over diameter ratios directly to percent of area. The procedure in using the nomograph is:

1. Start at the left-hand side of the nomograph with the trial diameter.

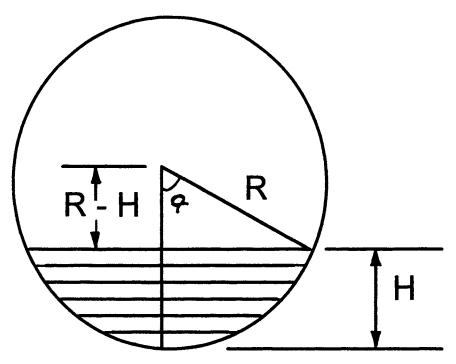


Figure 4-6. Area of the segment.

- 2. Align the internal diameter with the height of the vapor space and then with the rise of the minimum liquid level. Read the area occupied by these segments.
- 3. Subtract the sum of these segments from 100%. The difference is the area available for liquid holding time.

Chord length of segmental area, CL:

$$CL = 2(DH - H^2)^{0.5}$$
(4-28)

Partial volume of a horizontal cylinder:

$$V_{HC} = A_s L \tag{4-29}$$

Partial volume of dished heads:

 $V_{\rm DH} = 0.215483 {\rm H}^2 (1.5 {\rm D} - {\rm H})$ (4-30)

Partial volume of elliptical heads:

$$V_{\text{FLL}} = 0.52360 \,\text{H}^2 (1.5 \,\text{D} - \text{H}) \tag{4-31}$$

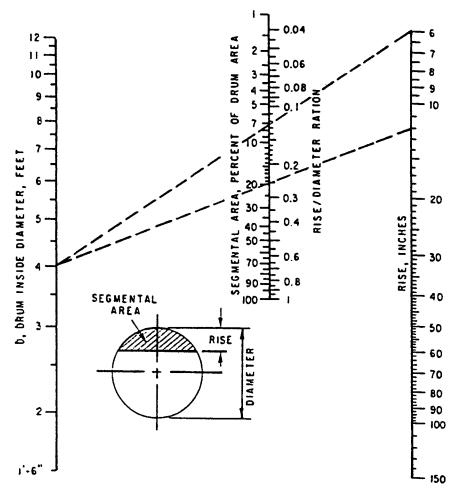


Figure 4-7. Nomograph to find segmental area available for liquid holding time.

Partial volume of hemispherical heads:

 $V_{\rm HS} = 2V_{\rm ELL} \tag{4-32}$

The total volume V_T :

$$\mathbf{V}_{\mathrm{T}} = \mathbf{V}_{\mathrm{HC}} + \mathbf{V}_{\mathrm{H}} \tag{4-33}$$

where V_{H} = head volume

Area =
$$\frac{\pi \cdot D^2}{4}$$
 (4-34)

Volume of a vertical tank:

$$V_{\rm VT} = 0.25\pi D^2 H \tag{4-35}$$

Conversions:

 $ft^3 = in^3/1728$ $gal = (in^3)(0.004329)$ $bbl = (in^3)(0.00010307)$

Nomenclature

Area = circle area. in^2 A_s = segmental area of a circle, in² CL = chord length of segmental area, inchbbl = volume in barrels d = vessel tank, or circle diameter, inch H = chord or liquid height, inchL = vessel length, inchR = radius of vessel or tank, inch V_{DH} = volume of dished head, in³ V_{FII} = volume of elliptical heads, in³ $V_{\rm H}$ = head volumes, in³ V_{HC} = volume of horizontal cylinder, in³ V_{HS} = volume of hemispherical heads, in³ V_{T} = total volume of horizontal vessel, in³ $V_{\rm VT}$ = volume of vertical tank, in³

PRELIMINARY VESSEL DESIGN

Columns and towers are the most essential aspects of the refinery system in the petrochemical or in the process chemical industries. They are divided into three separate functions: distillation, fractionation, and chemical operations. Stills are cylindrical chambers in which the application of heat to the charge stock changes from a liquid to a vapor. The vapor is then condensed in another vessel. Columns and towers are stills that increase the degree of separation that can be obtained during the distillation of crude oil. Fractionation towers are used for light end products. Generally, towers are large cylindrical vessels that have plates

(trays spaced inside the shell) to promote mixing of the downward flow of liquid with the upward flow of the vapors. These vessels require self-supporting, especially when they are exposed to high winds or to seismic risk.

Process conditions in the CPI often change, and a tower that is initially designed for vacuum operation may at a later stage be employed at pressure greater than atmospheric. Therefore, thin-walled vessels are often rated for both pressure and vacuum conditions. If the vessel is thick-walled and designed for high pressure service, no vacuum rating will be required. Tall vessels have one of a number of closures such as hemispherical, ellipsoidal, torispherical, conical, or flat ends.

The ellipsoidal shape is usual for closures of vessels that are six feet in diameter or greater. The hemispherical shape is preferred for vessels of a lesser diameter. The basic relationships for thin cylindrical shells under internal pressure assume that circumferential stress is dependent on the pressure and vessel diameter, but independent of the shell thickness.

That is:

$$f = \frac{P.D}{2t}$$
(4-36)

or

$$t = \frac{P.D}{2f}$$
(4-37)

Equation 4-36 is for membrane shells with a negligible thickness. As the pressure and the shell thickness increase, the stress distribution across the thickness is non-uniform. Therefore, some correction to the membrane theory is required. The modified code equation as given by the ASME Unfired Pressure Code, Section VIII, Division 1 is:

$$t = \frac{P.R}{S.E - 0.6P} + c$$
(4-38)

where c is the corrosion allowance

For a hemispherical head:

$$t_{\rm HH} = \frac{P.R}{2S.E - 0.2P}$$
(4-39)

For ellipsoidal dished head:

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$$t_{ELL} = \frac{P.D}{2S.E - 0.2P} + c$$
 (4-40)

For torispherical head:

$$t_{\text{TOR}} = \frac{0.885 \text{P.R}_{\text{c}}}{\text{S.E} - 0.1 \text{P}} + c$$
(4-41)

The weight of the vessel shell is:

$$W_{TS} = \frac{1}{12} (OD + t)(\pi)(SL) \left(\frac{t}{12}\right)(\rho_s)$$
(4-42)

The head blank diameter is:

$$BD = \frac{\{(OD)(H, Fac) + 2SF\}}{12}$$
(4-43)

Weight of head:

$$W_{TH} = 0.25(BD)^{2}(\pi) \left(\frac{t}{12}\right) (\rho_{s})(2)$$
(4-44)

The total weight:

$$W_{TOT} = W_{TS} + W_{TH}$$
(4-45)

The total weight excludes the nozzles, attachments, and vessel internals. Table 4-3 shows the head blank diameter factor for different types of head.

Nomenclature

- BD = head blank diameter, inch
 - c = corrosion allowance, inch
 - D = inside diameter of shell or head, inch
 - E = joint efficiency
- H.Fac = head blank diameter factor
 - OD = outside diameter of head, inch
 - P = internal design pressure, psig
 - R = inside radius, inch
 - R_c = crown radius of torispherical head, inch
 - $S = allowable stress, lb/in^2$

Head Type	OD/t	H.Fac
Torispherical	>50	1.09
*	30-50	1.11
	20-30	1.15
Ellipsoidal	>20	1.24
	10-20	1.30
Hemispherical	>30	1.60
1	18-30	1.65
	10-18	1.70

 Table 4-3

 Head Blank Diameter Factor (H.Fac)

SF = straight flange length, inch

SL =shell length, ft

t =shell thickness, ft

 t_{ELL} = ellipsoidal head thickness, inch

 t_{HH} = hemispherical head thickness, inch

 t_{TOR} = torispherical head thickness, inch

 W_{TH} = weight of head, lb

 W_{TS} = weight of vessel shell, lb

 W_{TOT} = total weight of vessel, lb

 ρ_s = density of vessel material, lb/ft³

CYCLONE DESIGN

Introduction

Cyclones are widely used for the separation and recovery of industrial dusts from air or process gases. Pollution and emission regulations have compelled designers to study the efficiency of cyclones. Cyclones are the principal type of gas-solids separators using centrifugal force. They are simple to construct, of low cost, and are made from a wide range of materials with an ability to operate at high temperatures and pressures. Cyclones are suitable for separating particles where agglomeration occurs. These types of equipment have been employed in the cement industry, and with coal gases for chemical feedstock, and gases from fluidized-bed combustion, as well as in the processing industry for the recovery of spray-dried products, and recently, in the oil industry for separating gas from liquid.

Cyclone reactors permit study of flow pattern and residence time distribution [6,7]. See for example, the studies by Coker [8,9] of synthetic detergent production with fast reaction. Reactor cyclones are widely used to separate a cracking catalyst from vaporized reaction products.

Reverse flow cyclones, in which the dust-laden gas stream enters the top section of the cylindrical body either tangentially or via an involute entry, are the most common design. The cylindrical body induces a spinning, vortexed flow pattern to the gas-dust mixture. Centrifugal force separates the dust from the gas stream; the dust travels to the walls of the cylinder and down the conical section to the dust outlet. The spinning gas also travels down the wall toward the apex of the cone, then reverses direction in an air-core and leaves the cyclone through the gas outlet tube at the top. This tube consists of a cylindrical sleeve and the vortex finder, whose lower end extends below the level of the feed port. Separation depends on particle settling velocities, which are governed by size, density, and shape.

Stairmand [10], Strauss [11], Koch and Licht [12] have given guidelines for designing cyclones. The effects of feed and cyclone parameters on the efficiency are complex, because many parameters are interdependent. Figure 4-3 shows the design dimensions of a cyclone and Table 4-4 gives the effects on cyclone performance in the important operating and design parameters.

Cyclone Design Procedure

The computation of a cyclone fractional or grade efficiency depends on cyclone parameters and flow characteristics of particle-laden gases. The procedure involves a series of equations containing exponential and logarithmic functions. Koch and Licht [12] described a cyclone using seven geometric ratios in terms of its diameter as:

$$\frac{a}{D_{c}}, \frac{b}{D_{c}}, \frac{D_{c}}{D_{c}}, \frac{S}{D_{c}}, \frac{h}{D_{c}}, \frac{H}{D_{c}}, \frac{B}{D_{c}}$$

They further stated that certain constraints are observed in achieving a sound design. These are:

$$a < S$$
 to prevent short circuiting

Variable	Effect
Pressure drop increases	Cut size (diameter of particles of which 50% are collected) decreases, flow rate increases; sharpness increases.
Solids content of feed increases	Cut size increases (large effect above 15–20% v/v).
$(\rho_p - \rho_f)$ increases	Cut size decreases.
Viscosity of liquid phase increases	Little effect below 10 mPas.
Cyclone diameter (D_c) increases	Cut size increases; pressure drop usually decreases.
Cyclone inlet (a) diameter increases	Gravitational force in cyclone decreases; cut size increases; capacity falls; pressure drop decreases.
Overflow diameter increases	Cut size increases; risk of coarse sizes appearing.
Underflow diameter increases	Brings excess fines from liquid phase into underflow.
Cyclone shape becomes longer	Decreases cut size; sharpens separation.

Table 4-4Effects of Variables on Cyclone Performance

$b < \frac{1}{2} (D_c - D_c)$) to avoid sudden contraction		
$S + l \le H$ to	keep the vortex inside the cyclone		
S < h			
h < H			
$\Delta P < 10$ in H_2O			
$v_i/v_s \le 1.35$	to prevent reentrainment		
$v_i^{}/v_s^{} \cong 1.25$	for optimum efficiency		

The Equations

Natural length, l: the distance below the gas outlet where the vortex turns.

$$1 = 2.3D_{e} \left(\frac{D_{c}^{2}}{ab}\right)^{1/3}$$
(4-46)

For 1 < (H - S), the cyclone volume at the natural length (excluding the core) is V_{nl} :

$$V_{nl} = \frac{\pi D_{c}^{2}}{4} (h - S) + \left(\frac{\pi D_{c}^{2}}{4}\right) \left(\frac{1 + S - h}{3}\right) \left(1 + \frac{d}{D_{c}} + \frac{d^{2}}{D_{c}^{2}}\right) - \frac{\pi D_{e}^{2} l}{4}$$
(4-47)

The diameter of a central core at a point of vortex turns, d:

$$d = D_{c} - (D_{c} - B) \left(\frac{S - l - h}{H - h} \right)$$
 (4-48)

For 1 > (H – S), the cyclone volume below the exit duct (excluding the core) $V_{\rm H}$:

$$V_{\rm H} = \frac{\pi D_{\rm c}^2}{4} (\rm h - S) + \left(\frac{\pi D_{\rm c}^2}{4}\right) \left(\frac{\rm H - h}{3}\right) \left(1 + \frac{\rm B}{\rm D_{\rm c}} + \frac{\rm B^2}{\rm D_{\rm c}^2}\right) - \frac{\pi D_{\rm e}^2}{4} (\rm H - S)$$
(4-49)

For a vortex exponent, n:

$$n = 1 - \left\{ 1 - \frac{(12D_c)^{0.14}}{2.5} \right\} \left\{ \frac{T + 460}{530} \right\}^{0.3}$$
(4-50)

The cyclone volume constant, K_c , using V_{nl} or V_{H} :

$$K_{c} = \frac{(2V_{s} + V_{nl,H})}{2D_{c}^{3}}$$
(4-51)

where

$$V_{s} = \left\{ \frac{\pi (S - a/2)(D_{c}^{2} - D_{e}^{2})}{4} \right\}$$
(4-52)

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The relaxation time (τ_i) for particle species i of diameter d_{pi} is

$$\tau_{i} = \frac{\rho_{p}(d_{pi})^{2}}{18\mu}$$
(4-53)

Cyclone configuration factor G: This is specified by the geometric ratios that describe the cyclone's shape. The cyclone configuration factor G is a function only of the configuration and is specified by the seven geometrical ratios that describe its shape. G is expressed as:

$$G = \frac{8K_{c}}{K_{a}^{2}K_{b}^{2}}$$
(4-54)

where

$$K_a = \frac{a}{D_c}, \quad K_b = \frac{b}{D_c}$$
(4-55)

Substituting the values of K_a , K_b , and K_c in Equation 4-54.

G = {2[
$$\pi(S - a/2)(D_c^2 - D_e^2)$$
] + 4V_{H,nl}} $\frac{D_c}{a^2b^2}$ (4-56)

The fractional or grade efficiency, η_i , can be expressed as:

$$\eta_{i} = 1 - \exp\left\{-2\left[\frac{G\tau_{i}Q}{D_{c}^{3}}(n+1)\right]^{0.5/(n+1)}\right\}$$
(4-57)

Saltation Velocity

Koch and Licht [12] expressed the saltation velocity as:

- The minimum fluid velocity necessary to prevent the settling out of solid particles carried in the stream.
- The necessary velocity that picks up deposited particles and transports them without settling.

Zenz [13] has shown that the velocity given by the latter differs from the former by a factor of 2 to 2.5. Kalen and Zenz [14] have applied the saltation concept to cyclone design by assuming:

- There is no slippage between fluid and particles. The cyclone inlet width is the effective pipe diameter for calculating saltation effects.
- Grain loading (dust concentration) is less than 10 grains/ft³.
- The diameter effect on the saltation velocity is proportional to the 0.4 power of the inlet width.

The saltation velocity, v_s , is dependent on cyclone dimensions, particle and fluid properties. v_s is expressed as:

$$v_{s} = 2.055\omega \left\{ \frac{b/D_{c}}{(1 - b/D_{c})^{1/3}} \right\} D_{c}^{0.067} v_{i}^{2/3}$$
 ft/s (4-58)

where

$$\omega = \left\{ \frac{4g\mu(\rho_{\rm p} - \rho_{\rm f})}{3\rho_{\rm f}^2} \right\}^{1/3}$$
(4-59)

Inlet velocity, v_i: ft/s

$$v_i = \frac{Q}{(ab)}$$
(4-60)

Kalen and Zenz have shown that maximum cyclone collection efficiency occurs at $v_i/v_s = 1.25$, and Zenz has found experimentally that fluid reentrainment occurs at $v_i/v_s = 1.36$.

Pressure Drop (ΔP)

Several attempts have been made to calculate the frictional loss or ΔP of a cyclone, although none has been very satisfactory. These are because assumptions made have not considered entrance compression, wall friction, and exit contraction, all of which have a major effect. Consequently, no general correlation of cyclone ΔP has been adopted. Pressure drop in a cyclone with collection efficiency is important in evaluating its cost. Correlations for the pressure drop have been empirical and are acceptable up to $\Delta P = 10$ in. H₂O. The pressure drop (ΔP) or the frictional loss is expressed in terms of the velocity head based on the cyclone inlet area. The frictional loss through cyclones is from 1 to 20 inlet velocity heads and depends on the geometric ratios. ΔP through a cyclone is given by

$$\Delta P = 0.003 \rho_f v_i^2 N_H \tag{4-61}$$

where

$$N_{\rm H} = K \left(\frac{ab}{D_{\rm e}^2} \right) \tag{4-61}$$

K = 16 for no inlet vane

K = 7.5 with a neutral inlet vane

 ΔP depends strongly on the inlet velocity and high velocities can cause both reentrainment and high pressure drop. However, entrainment can be reduced to a minimum, if the cyclone has a small base angle.

Trouble Shooting Cyclone Maloperations

In general, cyclones are used to separate particles from the gas stream, but recent developments have enabled cyclones to function as reactors. Some cyclone reactors can separate cracking catalyst from vaporized reaction products in the range of 950°F and 1,000°F, or can function as regenerators for flue gases between 1,250°F and 1,500°F. In both cases, the high particle velocities can cause rapid erosion of the cyclone material. This often results in poor performance of the cyclone. Other causes of poor cyclone performance are:

- Hole in cyclone body
- Cyclone volute plugged
- Dipleg unsealed
- Dipleg plugged
- Dipleg failure

Lieberman [15] has reviewed the causes of these maloperations, which often result in loss catalyst and reduced efficiency. A deficient cyclone reactor is identified by bottom sediment and water levels in the slurry oil product. For a regenerator cyclone, problems are visibly identified by the increased opacity of the regenerator flue gas or by reduced rates of spent catalyst withdrawal.

Cyclone Collection Efficiency

Many theories have been proposed to predict the performance of a cyclone, although no fundamental relationship has been accepted. Attempts have been made to predict the critical particle diameter, $(D_p)_{crit}$.

This is the size of the smallest particle that is theoretically separated from the gas stream with 50 percent efficiency. The critical particle diameter is defined by [16]

$$(\mathbf{D}_{p})_{crit} = \left[\frac{9\mu \mathbf{D}_{c}}{4\pi N_{t} \mathbf{v}_{i} (\rho - \rho_{g})}\right]^{0.5}$$
(4-62)

where N_t = effective number of turns made by the gas stream in the cyclone, and is defined by

$$N_{t} = (v_{i}) \left[0.1079 - 0.00077 v_{i} + 1.924(10^{-6}) v_{i}^{2} \right]$$
(4-63)

and $v_i = is$ the inlet linear velocity.

Figure 4-8 shows the percentage removal of particles in a cyclone as a function of the ratio of the particle to the critical diameter.

Cyclone Design Factor

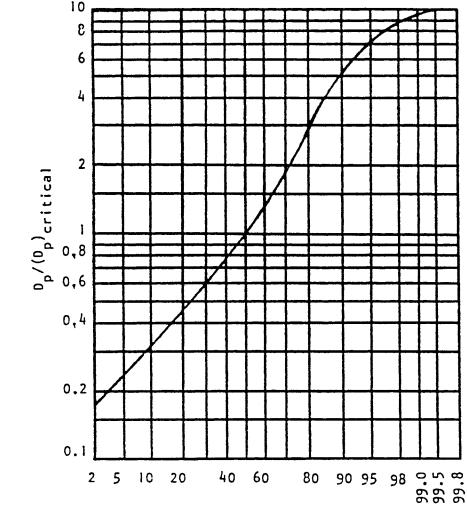
Cyclones are designed to meet specified ΔP limitations. The factor that controls the collection efficiency is the cyclone diameter, and a smaller diameter cyclone at a fixed ΔP will have a higher efficiency. Therefore, small diameter cyclones require a multiple of units in parallel for a given capacity. Reducing the gas outlet diameter results in an increased collection efficiency and ΔP . High-efficiency cyclones have cone lengths in the range of 1.6 to 3.0 times the cyclone diameters.

Collection efficiency increases as the gas throughput increases. Kalen and Zenz [14] reported that collection efficiency increases with increasing gas inlet velocity to a minimum tangential velocity. This reaches the point where the dust is re-entrained or not deposited because of saltation. Koch and Licht [12] showed that saltation velocity is consistent with cyclone inlet velocities in the range of 50 to 90ft/sec.

Cyclones offer the least expensive means of dust collection. They give low efficiency for collection of particles smaller than 5μ m. A high efficiency of 98 percent can be achieved on dusts with particle sizes of 0.1 to 0.2 μ m that are highly flocculated.

Cyclone Design Procedure

The following design procedures can be used to size a cyclone with a specified fluid flow rate and physical property data:



Percent removed

Figure 4-8. Percentage removal of particles as a function of particle diameter relative to the critical diameter.

- 1. Select either the high-efficiency or high-throughput design depending on the performance required.
- 2. Obtain an estimate of the particle size distribution of the solids in the stream to be treated.
- 3. Estimate the number of cyclones needed in parallel.
- 4. Calculate the cyclone diameter for an inlet velocity of 50 ft/sec.

- 5. Calculate the scale-up factor for the transposition of grade efficiency and particle size.
- 6. Calculate the cyclone performance and overall efficiency (recovery of solids). If unsatisfactory, try a smaller diameter.
- 7. Calculate the cyclone pressure drop and, if required, select a suitable blower.
- 8. Cost the system and optimize to make the best use of the pressure drop available or, if a blower is required, to give the lowest operating cost.

Nomenclature

- a = inlet height, ft
- b = inlet width, ft
- B = cyclone dust-outlet diameter, ft
- d = diameter of central core at a point where vortex turns, ft
- d_{pi} = diameter of particle in size range i, ft
- $D_c = cyclone diameter, ft$
- $D_e = cyclone gas-outlet diameter, ft$
- g = acceleration due to gravity, 32.2 ft/s^2
- G = cyclone configuration factor
- h = cylindrical height of cyclone, ft
- i = subscript denotes interval in particle size range

$$K_a = a/D_c$$

$$K_b = b/D_c$$

- $K_c = cyclone volume constant$
 - 1 = natural length (distance below gas outlet where vortex turns), ft
 - n = vortex exponent
- $N_{\rm H}$ = number of inlet velocity heads
- $\Delta \ddot{P}$ = pressure drop, in H₂O
 - Q = total gas flowrate, actual ft³/s
 - S = gas outlet length, ft
 - T = temperature, °F
 - $v_i = inlet velocity, ft/s$
- $v_s =$ saltation velocity, ft/s
- $V_{\rm H}$ = volume below exit duct (excluding core), ft³
- V_{nl} = volume at natural length (excluding core), ft³
- V_s = annular volume above exit duct to middle of entrance duct, ft³
- η_i = grade efficiency for particle size at midpoint of interval i, %
- $\mu =$ fluid viscosity, lb_m/ft.s
- $\rho_{\rm f}$ = fluid density, $lb_{\rm m}/ft^3$

 ρ_p = particle density, $lb_m/ft^3 \tau$ = relaxation time, s ϖ = Equation 4-59

GAS DRYER (DEHYDRATION) DESIGN

Liquid water and sometimes water vapor are removed from natural gas to prevent corrosion and formation of hydrates in transmission lines and to attain a water dew point requirement of the sales of gas. Many sweetening agents employ an aqueous solution for treating the gas. Therefore dehydrating the natural gas that normally follows the sweetening process involves:

- dehydration by refrigeration
- absorption by liquid desiccants
- adsorption by solid desiccants

Adsorption is a process that involves the transfer of a material from one phase to a surface where it is bound by intermolecular forces. The process involves the transfer from a gas or liquid to a solid surface. It could also involve the transfer from a gas to a liquid surface. The adsorbate is the material being concentrated on the surface and the material that it accumulates is defined as the adsorbent.

The oil and chemical industries use the adsorption process in the cleanup and purification of wastewater streams and for the dehydration of gases. The process is also used in gas purification involving the removal of sulfur dioxide from a stack gas. In addition, adsorption is employed to fractionate fluids that are difficult to separate by other separating methods. The amount of adsorbate that is collected on a unit of surface area is negligible. Therefore, porous desiccants (adsorbent) having a large internal surface area are used for industrial applications.

There are many solid desiccants that can adsorb water from natural gas. The common commercial desiccants are alumina, silica gel, and molecular sieves. The molecular sieves possess the highest water capacity, will give the lowest water dew points, and can be applied to sweeten dry gases and liquids. Figure 4-9 shows how desiccants can be used in dehydrators containing two or more towers. One of the towers operates using steam to adsorb water from the gas, while the other tower is regenerated and cooled. The regenerated gas is heated to a range of 450°F and 600°F, depending on the type of solid desiccant and the nature of service [17].

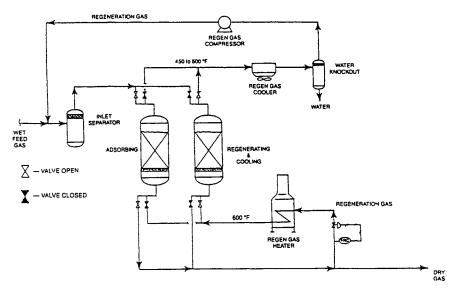


Figure 4-9. Solid desiccant dehydrator twin tower system. Courtesy: Gas Processors Suppliers Association *Engineering Data Book*, Tenth Edition.

The use of solid desiccant is limited to applications such as those with very low water dewpoint requirements, simultaneous control of water and hydrocarbon dewpoints, and in very sour gases. In cryogenic plant, solid-desiccant dehydration is much preferred over methanol injection to prevent hydrate and ice formation.

The basic design procedures for dehydrating saturated natural gas at a specified dew point are:

- 1. Determine the process conditions and the dryer process flow diagrams.
- 2. Select a drying cycle and calculate the water load.
- 3. Select the type of desiccant and compute the capacity and volume required.
- 4. Size the dryer and check for the pressure drop.
- 5. Calculate the desiccant reactivation heating and cooling requirements.

Here, design procedures are presented for sizing a solid desiccant dryer in removing moisture from gas streams. The process involves the following calculations.

- 1. the water pickup
- 2. desiccant volume

- 3. desiccant bed
- 4. gas velocity through the bed
- 5. vessel weight
- 6. dryer regeneration requirements

The design is based upon the following assumptions [2]:

- 1. The temperature difference between the heater outlet temperature and the peak vessel outlet temperature is 50°F.
- 2. The average bed temperature is based on 75 percent of the bed at heater outlet temperature and 25 percent at the peak vessel outlet temperature Equation 4-73.
- 3. Specific heats: steel, 0.12 Btu/lb °F, and desiccant 0.25 Btu/lb °F.
- 4. Heat of water desorption 1400 Btu/lb H₂O adsorbed.
- 5. Flat heads used on vessel ends; steel density is 480 lb/ft³.
- 6. Total vessel weight increased by 10 percent for supports.
- 7. Heat losses to the dryer during heating period calculated at 5 percent.

The Equations

The equations used for calculating these requirements are: The total water adsorbed, lb

$$H_{2}O \text{ load} = \frac{(\text{Flow})(\text{Cycle})(\text{H}_{2}O \text{ Cont})}{24}$$
$$= \frac{\left(\frac{\text{MMstd ft}^{3}}{\text{day}}\right)(h)\left(\frac{\text{lb}}{\text{MMstd ft}^{3}}\right)}{(h/\text{day})}$$
(4-64)

Desiccant volume

$$V_{\text{DES}} = \frac{(H_2 O \text{ load})(100)}{(\text{pickup})(\rho_D)}$$
(4-65)

The calculated area based on desiccant volume

$$A_{\text{DES}} = \frac{V_{\text{DES}}}{L}$$
(4-66)

The bed diameter

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$$D_{BED} = \left(\frac{4A_{DES}}{\pi}\right)^{0.5}$$
(4-67)

actual desiccant bed area

$$A = \frac{\pi D_{BED}^2}{4}$$
(4-68)

The actual gas flowrate at flowing conditions, ft3/min

$$Q = (Flow) \left(\frac{14.7}{P + 14.7} \right) \left(\frac{T}{520} \right) \left(\frac{1}{1440} \right) (Z)$$

= (Flow) $\left(\frac{T}{P + 14.7} \right) (Z) (19.6314)$ (4-69)

Superficial gas velocity, ft/min

$$V_{GAS} = \frac{Q}{A}$$
(4-70)

Superficial gas velocities for adequate contact time should range between 30 and 60ft/min

Dryer shell thickness, inch

$$t = \frac{PR}{SE - 0.6P}$$
(4-71)

Vessel weight

$$W_{VES} = 480 \pi D \left(\frac{t}{12}\right) (L + h + R)(1.10)$$
(4-72)

Add an extra 2 ft (h = 2) of straight side (i.e., for 6 in. of 3/8-in. inert balls above and below the desiccant bed and 1 ft for an inlet-gas distributor).

Temperature of desiccant bed, °F

$$T_{BED} = 0.75(HT) + 0.25(HT - 50)$$
(4-73)

Total heat required to regenerate the dryer (ideal), Btu

$$H_{TOT} = \{ (W_{VES})(0.12)(T_{BED} - T) + (W_{D})(0.25)(T_{BED} - T) + (H_2O \log)(1400) \} (1.05)$$
(4-74)

where

$$W_{\rm D} = (\rho_{\rm D})(V_{\rm DES})$$

loss factor for non-steady state heating, F

$$F = \ln\left(\frac{HT - T}{50}\right) \tag{4-75}$$

Heat added to regeneration gas to regenerate the desiccant bed

$$H_{REG} = H_{TOT}.F$$
(4-76)

The total regeneration gas requirement, lb

$$H_{GAS} = \frac{H_{REG}}{H_1 - H_2}$$
(4-77)

Standard cubic feet of regeneration gas

$$SCF = \frac{H_{GAS}(380)}{MW}$$
(4-78)

Pressure Drop (ΔP)

The pressure drop is estimated from the desiccant manufacturer's data and correlations. In addition, ΔP can be calculated by Wunder's [18] graphical correlation of gas superficial velocity against ΔP in ft water per ft bed. Wunder used a factor of 1.6 for a fouled bed. ΔP through a new desiccant bed is initially low. After a short time in service, ΔP rises because of bed settling and then slowly increases over the active life of the desiccant because of more settling and some attrition. Figure 4-10 shows ΔP for an 8-inch silica gel desiccant. These data are based on air. For other gases, the given values should be multiplied by (MW gas/MW air)^{0.9}. The pressure drop for a clean bed is given by:

$$\Delta \mathbf{P} = \left(\frac{\mathrm{ft}\,\mathrm{H}_{2}\mathrm{O}}{\mathrm{ft}\,\mathrm{bed}}\right) \left(\frac{0.4335\,\mathrm{psi}}{\mathrm{ft}\,\mathrm{H}_{2}\mathrm{O}}\right) \left(\frac{\mathrm{MWgas}}{\mathrm{MWair}}\right)^{0.9} \tag{4-79}$$

Fouled bed:

$$\Delta P = \text{clean bed } \Delta P \times 1.6 \tag{4-80}$$

Desiccant attrition due to high gas velocities is controlled by proper bed design. The superficial gas velocity is used to estimate Alcoa's

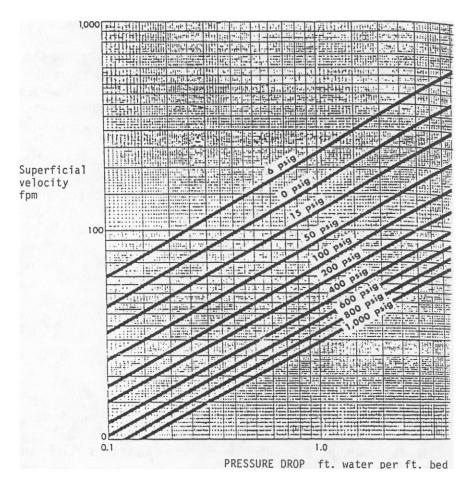


Figure 4-10. Pressure drop for an 8 mesh silica gel desiccant (by permission Oil & Gas Journal).

momemtum downflow velocities [18]. However, desiccant attrition should not be a problem when the momentum number is equal to or less than 30,000. Although, this number is based on granular alumina, it can be used for silica gel, which has crushing characteristics similar to alumina. The momentum is expressed as:

Momentum =
$$(V_{GAS})(MW)\left(\frac{P+14.7}{14.7}\right) \le 30,000$$
 (4-81)

where V_{GAS} = superficial gas velocity, ft/min MW = molecular weight of gas P = system pressure in psig

Desiccant Reactivation

The desiccant bed is reactivated at as high a temperature as permissible to start optimum capacity and minimum dew points. In natural gas drying service, the normal reactivation temperature for silica gel is between 350°F and 400°F. Because the feed gas contains high pentanes-plus materials, Wunder [18] suggested that reactivation can be carried out at 450°F to desorb the heavy hydrocarbons.

At a temperature above 450°F, the attrition rate for silica gel is high. However, occasionally at 600°F, reactivation will restore some desiccant's initial activity thus extending the life of the silica gel. Therefore, the reactivation procedure is designed to periodically heat the desiccant to 600°F [18].

Nomenclature

A = actual desiccant bed area, ft^2

 A_{DES} = calculated area based on desiccant volume, ft²

Cycle = drying cycle, h

- D_{BED} = calculated bed diameter, ft
 - E = joint efficiency
 - F = loss factor for non-steady state heating
- Flow = gas flow rate, 10^6 std ft³/day
 - h = additional shell height for desiccant supports and distributor, ft
- H_1 , H_2 = enthalpy of regeneration gas before and after regeneration heater, Btu/lb
 - H_{GAS} = total regeneration gas requirement, lb
 - H_{REG} = heat added to regeneration gas to regenerate desiccant bed
 - H_{TOT} = heat required to regenerate dryer (ideal), Btu
 - HT = heater temperature, °F
- H_2O load = total water adsorbed, lb
- H_2O Cont = water content of gas, $lb/10^6$ std ft³
 - L = desiccant bed length, ft
 - MW = molecular weight of regeneration gas
 - P = dryer operating pressure, psig
- PICKUP = useful design capacity of desiccant, %
 - Q = actual gas flow rate at flowing conditions, ft^3/min
 - R = radius of desiccant bed, ft
 - $S = allowable stress, lb/in^2$
 - SCF = standard cubic feet of regeneration gas

t = dryer shell thickness, inch

T = gas flowing temperature, $^{\circ}R$

TB = temperature of desiccant bed, °F

 V_{DES} = desiccant volume, ft³

 V_{GAS} = superficial gas velocity, ft/min

 W_D = desiccant weight, lb

WS = steel weight of shell and heads, lb

 W_{VES} = vessel weight, lb

Z = compressibility factor

 $\rho_{\rm D}$ = desiccant density, lb/ft3

PROBLEMS AND SOLUTIONS

Problem 4-1

Size a vertical separator under the following conditions:

WL = 5000 lb/h	WV = 37,000 lb/h
$\rho_{\rm L} = 61.87 \ \rm lb/ft^3$	$\rho_v = 0.374 \text{ lb/ft}^3$

The liquid surge time is five minutes.

Size a horizontal separator for the following conditions:

WL = 56,150 lb/h	WV = 40,000 lb/h
$\rho_{\rm L} = 60.0 \rm lb/ft^3$	$\rho_{\rm v} = 1.47 \ {\rm lb/ft^3}$

The liquid surge time is six minutes.

Use L/D = 3.0

Solution

The computer program PROG41 sizes both the vertical and horizontal separators. Table 4-5 shows the input data and computer output for both separators. From the computed output, the diameter of the vertical separator with the given flow data and physical properties is 2.88 ft, the liquid height is 1.03 ft and the vessel volume is 6.74 ft³. The computer output for the horizontal separator gives its diameter as 4.13 ft, the vessel length is 12.39 ft, and volume is 165.82 ft³.

Figure 4-7 gives a nomograph of the vapor rise in horizontal vessels. A check of the vapor area with 20 percent of the total cross-sectional

 Table 4-5

 Input Data and Computer Output for

 Vertical and Horizontal Separators

DATA41.DAT	
1 5000.0 37000.0 61.87 0.374 5.0 2 56150.0 40000.0 60.0 1.47 6.0 3.0	
VERTICAL SEPARATOR SIZING	

LIQUID FLOWRATE, 1b/hr.:	5000.
VAPOR FLOWRATE, 1b/hr.:	37000.
LIQUID DENSITY, 1b/ft^3.:	61.870
VAPOR DENSITY, 1b/ft^3.:	.374
LIQUID VOLUMETRIC RATE, ft^3/sec.:	92.1232
VAPOR VOLUMETRIC RATE, ft^3/sec.: LIQUID SURGE TIME, min.:	.0224 5.0
SEPARATION FACTOR:	.0105
VAPOR VELOCITY FACTOR:	.329
MAXIMUM VAPOR VELOCITY, ft/s.:	4.213
VESSEL DIAMETER, ft:	2.882
MINIMUM VAPOR-LIQUID NOZZLE VELOCITY, ft/s.:	27.481
MAXIMUM VAPOR-LIQUID NOZZLE VELOCITY, ft/s.:	153.539
REQUIRED VESSEL VOLUME, ft^3:	6.735
LIQUID HEIGHT, ft.:	1.032
HORIZONTAL SEPARATOR SIZING ************************************	**************************************
LIQUID DENSITY, 1b/ft^3.:	60.000
VAPOR DENSITY, 1b/ft^3.:	1.470
LIQUID VOLUMETRIC RATE, ft^3/sec.:	.2600
VAPOR VOLUMETRIC RATE, ft^3/sec.:	7,5586
LIQUID SURGE TIME, min.:	6.000
LENGTH TO DIAMETER RATIO (L/D):	3.0
SEPARATION FACTOR:	.220
VAPOR VELOCITY FACTOR:	. 447
VAPOR VOLUMETRIC RATE, ft^3/s.:	7.559
MAXIMUM VAPOR VELOCITY, ft/s.:	2.823
MAXIMUM VAPOR VELOCITÝ, ft/s.: VESSEL DIAMETER, ft.:	2.823 4.129
MAXIMUM VAPOR VELOCITÝ, ft/s.: VESSEL DIAMETER, ft.: VESSEL LENGTH, ft.:	2.823 4.129 12.386
MAXIMUM VAPOR VELOCITÝ, ft/s.: VESSEL DIAMETER, ft.:	2.823 4.129

area and the computed diameter of 4.13 ft shows that the vapor rise is approximately 12 inches.

Problem 4-2

Estimate the liquid volume in a horizontal storage tank that has a diameter of 120 inches, a length of 200 inches, and contains 40 inches of liquid. Calculate the liquid volume for a vertical tank and for tanks with either flat heads, dished heads, elliptical and hemispherical.

Solution

The computer program PROG42 calculates the liquid volume in a horizontal tank and for tanks with flat, dished, elliptical and hemispherical heads. Table 4-6 shows the input data and computer output for a vertical tank and tanks with flat, dished, elliptical and hemispherical heads.

Problem 4-3

Estimate the weight of a pressure vessel with hemispherical heads. The vessel is 2 ft 6 in ID \times 10 ft – 0 in T/T and has a design pressure of 1000 psig.

2SF = 4 in	$E_{shell} = 0.85$
c = 0.125 in	$E_{head} = 1.0$
$\rho_s = 489.024 \text{ lb/ft}^3$	$S = 17,500 \text{ lb/in}^2$

Solution

The computer program PROG43 determines the vessel weight of the pressure vessel with hemispherical, elliptical, and torispherical heads. Table 4-7 gives input data and computer output for 2 ft 6 in.-diameter vessel. The output gives the shell and head thicknesses, the weights of the shell and head, and the total weight of the vessel with hemispherical, elliptical and torispherical heads.

Problem 4-4

Determine the efficiency and pressure drop (ΔP) based on cyclone dimensions and the gas flow rate at 516.7 ft³/s. and a density of 0.075 lb/ft³ containing particles with density of 62.43 lb/ft³.

The dimensions and data required are:

Cyclone gas inlet height, ft	4.5
Cyclone gas inlet width, ft	1.896
Cyclone gas outlet length, ft	3.448
Cyclone gas outlet diameter, ft	3.792
Cyclone overall height, ft	26.333
Cyclone cylindrical height, ft	8.552
Cyclone diameter, ft	6.333

(text continued on page 296)

Table 4-6

Input Data and Computer Outputs for Tank Volume with Flat, Dished, Elliptical and Hemispherical Heads, and Vertical Tank Volume

DATA4	2.DAT	1		
1				
120. 2	40.	200.		
120. 3	40.	200.		
120. 4	40.	200.		
120. 5	40.	200.		
120.	40.	200.		
				TANK VOLUME WITH FLAT HEADS

		DEPTH	R, in.:	120.000 40.000
		ENGTH,		200.000
Ť	ANK V	OLUME.	in.^3:	.660017E+06
т	'ANK V	OLUME.	ft.^3:	.381954E+03
т	ANK V	OLUME,	gal:	.285721E+04
T	ANK V	OLUME,	bb1:	.680279E+02
				TANK VOLUME WITH DISHED HEADS
			R, in.:	120.000
		DEPTH		40.000
		ENGTH,		200.000
		OLUME,		.662430E+06
Ť	ANK V	OLUME.	ft.^3:	.383351E+03
Ť	ANK V	OLUME,	gal:	.286766E+04
		OLUME,		.682767E+02
				TANK VOLUME WITH ELLIPTICAL HEADS
			R, in.:	120.000
		DEPTH		40.000
		ENGTH,		200.000
T	ANK V	OLUME,	in.^3:	.894589E+06
т	ANK V	OLUME.	ft.^3:	.517702E+03
т	ANK V	OLUME,	gal:	.387268E+04
T 	ANK V	OLUME,	bb1:	.922053E+02
*****	*****	******	******	TANK VOLUME WITH HEMISPHERICAL HEADS
			R, in.:	120.000
		DEPTH		40.000
т	ANK L	ENGTH,	in.:	200.000
т	ANK W	OLIME	in ^3.	.112916E+07
Т	ANK V	OLUME,	ft.^3:	.653450E+03
г	ANK V	OLUME,	gal:	.488814E+04
T 	ANK V	OLUME,	bbl:	.116383E+03
*****	*****	******	******	VERTICAL TANK VOLUME
			R, in.:	120.000
		DEPTH		40.000
		ENGTH,		200.000
-		,		2001000

Table 4-6 (continued)

-			
TANK	VOLUME,	in.^3:	.452389E+06
TANK	VOLUME,	ft.^3:	.261799E+03
TANK	VOLUME,	gal:	.195839E+04
TANK	VOLUME,	bbl:	.466278E+02

Table 4-7

Input Data and Computer Outputs for Vessel Design with Hemispherical, Ellipsoidal and Torispherical Heads

DATA4	3.DAT				
1					
-	10.0	1000.0			
2.0	0.125	489.024			
0.85	1.0	17500.			
2					
30.0	10.0	1000.0 489.024			
	1.0	17500.0			
3					
30.0	10.0	1000.0 489.024			
		489.024 17500.0 2.0			
0.85	1.0	17500.0 2.0			
			HEMISPHERICAL HEADS		
		DIAMETER, in.:	30.000		
		NGHT, ft.:	10.000		
		DESIGN PRESSURE, psig:	1000.000		
		FLANGE LENGTH, in.:	2.000		
		N ALLOWANCE, in.:	.125		
		OF VESSEL MATERIAL, 1b/ft^3.:	489.024		
		FICIENCY OF SHELL:	.850		
		FICIENCY OF HEAD:	1.000		
		E STRESS, lb/in^2.:	17500.000		
		ED SHELL THICKNESS, in.:	1.176		
C	ALCULAT	ED HEAD THICKNESS, in.:	.431		
		F VESSEL SHELL, 1b:	4205.77		
		F VESSEL HEAD, 1b:	595.80		
	TOTAL VE	SSEL WEIGHT, 1b:	4801.58		
		VESSEL DESIGN WITH	ELLIPSOTDAL HEADS		
*****	******		*****		
I	NTERNAL	DIAMETER, in.:	30.000		
S	SHELL LE	NGTH, ft.:	10.000		
I	INTERNAL	DESIGN PRESSURE, psig.:	1000.000		
S	STRAIGHT	FLANGE LENGTH, in.:	2.000		
		N ALLOWANCE, in.:	.125		
		OF VESSEL MATERIAL, 1b/ft^3.:	489.024		
	JOINT EF	.850			
	JOINT EFFICIENCY OF HEAD: 1.000				
A	ALLOWABLE STRESS, lb/in^2.: 17500.000				
		ED SHELL THICKNESS, in.:	1.176		
		ED HEAD THICKNESS, in.:	.987		
		F VESSEL SHELL, 1b:	4205.77		
		F VESSEL HEAD, 1b:	853.98		
	OTAL VE	SSEL WEIGHT, 1b:	5059.75		

VESSEL DESIGN WITH TORISPHERICAL HEADS

(table continued on next page)

-

Table 4-7 (continued)

INTERNAL DIAMETER, in.:	30.000
SHELL LENGTH, ft.:	10.000
INTERNAL DESIGN PRESSURE, psig.:	1000.000
STRAIGHT FLANGE LENGTH, in.:	2.000
CORROSION ALLOWANCE, in.:	.125
DENSITY OF VESSEL MATERIAL, 1b/ft^3.:	489.024
JOINT EFFICIENCY OF SHELL:	.850
JOINT EFFICIENCY OF HEAD:	1.000
ALLOWABLE STREES, 1b/in.^2:	17500.000
CALCULATED SHELL THICKNESS, in.:	1.176
CALCULATED HEAD THICKNESS, ft.:	. 227
WEIGHT OF VESSEL SHELL, 1b:	4205.77
WEIGHT OF VESSEL HEAD, 1b:	155.37
TOTAL VESSEL WEIGHT, 1b:	4361.15

(text continued from page 293)

2.533
516.7
0.075
$1.28 imes 10^{-5}$
110
3.281×10^{-5}

Solution

The computer program PROG44 calculates the pressure drop and the efficiency of the cyclone using the cyclone geometry, fluid and particle specifications. Table 4-8 gives the input data and computer results with an efficiency of 63.305%, and a pressure drop of 7.834 in H₂O. The calculated ratio of the particle diameter relative to the critical diameter is 0.51. From Figure 4-8, the percentage removal of the particle is 23%.

Problem 4-5

Size a dryer to operate under the following conditions as shown in Table 4-9 using silica gel as the adsorbent.

Solution

The computer program PROG45 calculates the size of the dryer using silica gel, and Table 4-10 illustrates the input data and computer output. The total regeneration flow is 44649.0 lb and the momentum is 30,584. The heat requirement for dryer regeneration is 5.6×10^8 Btu. The calculated superficial gas velocity is 34.95ft/min.

 Table 4-8

 Input Data and Computer Output for Cyclone Design

DATA44.DAT

26.333	8.552	6.333	2.533
4.5	1.896	3.448	3.792
516.7	62.43	0.075	0.0000128
110.0	0.0000	3281	

CYCLONE DESIGN	
CYCLONE OVERALL HEIGHT, ft:	26.333
CYCLONE CYLINDRICAL HEIGHT, ft.:	8.552
CYCLONE DIAMETER, ft.:	
· · · · · · · · · · · · · · · · · · ·	6.333
CYCLONE DUST-OUTLET DIAMETER, ft:	2.533
CYCLONE GAS INLET LENGTH, ft:	4,500
CYCLONE GAS INLET WIDTH, ft:	1.896
CYCLONE GAS OUTLET LENGTH, ft:	3.448
CYCLONE GAS OUTLET DIAMETR, ft:	3.792
GAS RATE, ft^3/s.:	516,70
PARTICLE DENSITY, 1b/ft^3:	62.430
GAS DENSITY, 1b/ft^3:	.075
GAS VISCOSITY, lb/ft.s.:	.128000E-04
GAS TEMPERATURE, OF:	110,000
PARTICLE SIZE (EQUIVALENT DIA.), ft.:	.328100E-04
VORTEX EXPONENT:	.7276
RELAXATION TIME, s:	.291691E-03
INLET FLUID VELOCITY, ft/s.:	60,560
SALTATION VELOCITY, ft/s.:	22.108
CYCLONE CONFIGURATION FACTOR:	89.728
CYCLONE PRESSURE DROP, in H2O:	7.834
RATIO OF THE INLET TO THE SALTATION VELOCITIES:	2.739
CYCLONE EFFICIENCY, %:	63,305
EFFECTIVE NUMBER OF TURNS:	3.71
CRITICAL PARTICLE DIAMETER, ft.:	.643700E-04
CRITICAL PARTICLE DIAMETER, IL.: CRITICAL PARTICLE DIAMETER, microns:	
RATIO OF PARTICLE DIAMETER, MICFORS: RATIO OF PARTICLE DIA. TO CRITICAL DIA.:	.196200E+02
RATIO OF PARTICLE DIA. TO CRITICAL DIA.:	.51

Table 4-9 Dryer Parameters

Flow = 65 MM std ft ³ /day	Desiccant H_2O cap = 8%
P = 700 psig	Desiccant density = 50 lb/ft^3
$T = 80^{\circ}F$	Desiccant bed length = 15ft
Z = 0.9228	Heater outlet = 500° F
H_2O Cont = 47 lb/MM std ft ³	MW = 18
$S = 15,500 \text{ lb/in}^2$	$H_1 = 490 \text{ Btu/lb}$
E = 1.0	$H_2 = 225 \text{ Btu/lb}$
Cycle = 12h	h = 2.0ft

From Figure 4-10,

Superficial gas velocity = 35 ft/min

Desiccant bed length = 15.0 ft

System pressure = 700 psig

The clean bed ΔP using Equation 4-79 is

$$\Delta P = 0.87 \frac{\text{ft H}_2\text{O}}{\text{ft bed}} \left(\frac{0.4335 \text{psi}}{\text{ft H}_2\text{O}} \right) (15.0 \text{ ft bed}) \left(\frac{18}{29} \right)^{0.9}$$

= 3.68psi

 Table 4-10

 Input Data and Computer Output for Gas Dryer Design

DATA45.	DAT		
65.0			
0.9228			
15500.0	1.0	8.0	
50.0	15.0	500.0	
18.0	490.0	225.0	
		GAS DRYER DESIGN	

		ATE, 10^6 std. ft^3/day:	65.000
		ATING PRESSURE, psig.:	700.000
		G TEMPERATURE, OF:	80.0
		SSIBILITY FACTOR: LE, hr.:	.9228 12.0
			47.
			15500.0
			1.0
		8.0	
		DENSITY, 1b/ft^3:	50.0
HEA	TER OUT	LET TEMPERATURE, OF:	500.0
		WEIGHT OF REGENERATION GAS:	18.0
		F GAS BEFORE REGENERATION HEATER, Btu/lb:	
		F GAS AFTER REGENERATION HEATER, Btu/lb:	225.0
		R ADSORBED, 1b/h.:	1527.500
		VOLUME, ft^3.:	381.8750
		CROSS-SECTIONAL AREA, ft^2.:	25.4583 5.6934
	RADIUS	ER, ft.:	34.1603
	LENGTH		15.00
		FLOW RATE, ft^3/min.:	889.697
		L GAS VELOCITY, ft./min.:	34.947
		L THICKNESS, in.:	1,5857
		GHT, 1b.:	24767.
		WEIGHT, 1b.:	19094.
TEM	PERATUR	E OF DESICCANT BED, oF:	487,500
TOT	AL HEAT	REQUIRED TO REGENERATE DRYER, Btu:	.555954E+07
LOS	S FACTO	R:	2.1282
HEAT	r added	TO REGENERATION GAS:	.118320E+08
TOT	AL REGE	NERATION GAS REQUIREMENT, 1b:	.446490E+05
STA	NDARD C	NERATION GAS REQUIREMENT, 15: UBIC FEET OF REGENERATION GAS, std.ft^3:	
MOMI	ENTUM:		.305838E+05

Fouled $\Delta P = 3.68 \times 1.6$

= 5.89 psi

The actual regeneration flow rate (lb/h) will depend upon the length of the heating cycle used.

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PROGRAM PROG41

с	********	***************************************
с	THIS PROGRAM	M USES THE WELL-KNOWN SOUDERS AND BROWN CORRELATION
с	COUPLED WIT	H LIQUID SURGE CAPACITY TO SIZE LIQUID-VAPOR SEPARATORS.
с	THE PROGRAM	SIZES BOTH VERTICAL AND HORIZONTAL SEPARATORS
с	*********	******************
c	HL =	LIQUID HEIGHT, ft.
č	SFac(SF)=	SEPARATION FACTOR
č	KV =	VAPOR VELOCITY FACTOR FOR VERTICAL SEPARATORS
c	VM =	MAXIMUM VAPOR VELOCITY, ft/s
c	WL =	LIQUID FLOW RATE, 1b/hr.
c	WV =	
c		VAPOR FLOW RATE, 1b/hr.
	DENL =	LIQUID DENSITY, 1b/ft^3.
С	DENV =	VAPOR DENSITY, 1b/ft^3
с	QV =	VAPOR VOLUMETRIC FLOW, ft^3/s
c	D =	VESSEL DIAMETER, ft
с	λ =	CROSS-SECTIONAL AREA OF VERTICAL SEPARATOR, ft^2
с	AV =	MINIMUM CROSS-SECTIONAL AREA OF VERTICAL SEPARATOR, ft^2
с	QL =	LIQUID VOLUMETRIC FLOW, ft^3/s
с	т =	LIQUID SURGE TIME, min.
с	KH =	VAPOR VELOCITY FACTOR FOR HORIZONTAL SEPARATORS
с	λT =	TOTAL CROSS-SECTIONAL AREA OF HORIZONTAL VESSEL, ft^2
с	L =	HORIZONTAL VESSEL LENGTH, ft
с	AL =	CROSS-SECTIONAL AREA OF HORIZONTAL VESSEL OCCUPIED BY
c		LIQUID PHASE, ft^2
c	X =	ln(S.Fac)
č		**********
č		
č		
•	REAL KH, KV COMMON B, D	1, L , E, F, G, PI
	COMMON/DATA COMMON/DATA COMMON/DATA	1/ WL1, WV1, DENL1, DENV1, T1 2/ SF1, KV1, VMAX1, D1 3/ UMAX, UMIN, QL1, QV1, V, HL 4/ WL2, WV2, DENL2, DENV2, RATIO 5/ SF2, KH, QL2, QV2, VMAX2, D2, L, VESVOL, T2, CALT2
		3, FILE='DATA41.DAT', STATUS='OLD', ERR=18) 1, FILE='PRN')
с	PI = 3.1415	927
c	CONSTANTS F	OR VAPOR VELOCITY FACTOR
	B = -1.8774 D = -0.8145 E = -0.1870 F = -0.0145 G = -0.0010	80 74 23
5	READ (3, *,	ERR=19) I
	GO TO (11,	22),I
c	READ LIQUID	FLOW RATE WII. 1b/br

с с с с	READ VAPOR FLOW RATE, WV1: lb/hr. READ LIQUID DENSITY, DENL1: lb/ft^3. READ VAPOR DENSITY, DENV1: lb/ft^3. READ LIQUID SURGE TIME, T1: min.
11	READ (3, *, ERR=19) WL1, WV1 READ (3, *, ERR=19) DENL1, DENV1 READ (3, *, ERR=19) T1
	GO TO 33
000000	READ LIQUID FLOW RATE, WL2: lb/hr. READ VAPOR FLOW RATE, WV2: lb/hr. READ LIQUID DENSITY, DENL2: lb/ft^3. READ VAPOR DENSITY, DENV2: lb/ft^3. READ LIQUID SURGE TIME, T2: min. READ LENGTH-TO-DIAMETER RATIO, RATIO.
22	READ (3, *, ERR=19) WL2, WV2 READ (3, *, ERR=19) DENL2, DENV2, T2 READ (3, *, ERR=19) RATIO
	GO TO 44
18 20	WRITE (*, 20) Format (6X, 'Data file does not exist') Go to 999
19 21	WRITE (*, 21) Format (6x, 'Error message in the data value') Go to 999
с	COMPUTE THE SIZING OF THE VERTICAL SEPARATOR
33	CALL VERT
с	OUTPUT THE RESULTS OF THE VERTICAL SEPARATOR
	CALL OUTV
	GO TO 5
с	COMPUTE THE SIZING OF THE HORIZONTAL SEPARATOR
44	CALL HORIZ
с	OUTPUT THE RESULTS OF THE HORIZONTAL SEPARATOR CALL OUTH
	GO TO 999
	CLOSE (3, STATUS='KEEP') CLOSE (1)
999	STOP END
c c	**************************************

```
c
c
        SUBROUTINE VERT
       REAL KV1
       COMMON B, D, E, F, G, PI
COMMON/DATA1/ WL1, WV1, DENL1, DENV1, T1
COMMON/DATA2/ SF1, KV1, VMAX1, D1
COMMON/DATA3/ UMAX, UMIN, QL1, QV1, V, HL
с
        CALCULATE THE SEPARATION FACTOR
        SF1 = (WL1/WV1)*SQRT(DENV1/DENL1)
        X1 = ALOG(SF1)
        VAL1 = B+(D*X1)+(E*(X1**2))+(F*(X1**3))+(G*(X1**4))
С
        CALCULATE THE VAPOR VELOCITY FACTOR, ft./s.
        KV1 = EXP(VAL1)
с
        CALCULATE THE MAXIMUM VAPOR VELOCITY, ft./s.
        VMAX1 = KV1*SQRT((DENL1-DENV1)/DENV1)
С
       CALCULATE THE VOLUMETRIC RATE, ft^3/sec.
        QV1 = WV1/(DENV1*3600.0)
        AV1 = QV1/VMAX1
с
       CALCULATE THE DIAMETER OF THE VESSEL, ft.
        D1 = SQRT((4.0 * AV1)/PI)
        AREA = (PI*D1**2)/4.0
с
с
        CALCULATE THE VAPOR-LIQUID INLET NOZZLE VELOCITIES.
        DENM = (WL1+WV1)/((WL1/DENL1)+(WV1/DENV1))
        UMAX = 100.0/(DENM**0.5)
        UMIN = 60.0/(DENM**0.5)
C
        CALCULATE THE LIQUID VOLUMETRIC RATE, ft^3/sec.
        QL1 = WL1/(DENL1*3600.)
       CALCULATE THE REQUIRED VESSEL VOLUME, ft^3.
С
        V = (QL1*T1*60.0)
C
       CALCULATE THE LIQUID HEIGHT, ft.
       HL = V/AREA
       RETURN
        END
С
С
        С
       THIS PROGRAM SIZES HORIZONTAL LIQUID-VAPOR SEPARATOR
        ******
с
        SUBROUTINE HORIZ
        REAL L, KH, KV2
       COMMON / B, D, E, F, G, PI
COMMON/DATA4/ WL2, WV2, DENL2, DENV2, RATIO
COMMON/DATA5/ SF2, KH, QL2, QV2, VMAX2, D2, L, VESVOL, T2, CALT2
```

CALCULATE THE VAPOR-LIQUID SEPARATION FACTOR с SF2 = (WL2/WV2)*SQRT(DENV2/DENL2) X2 = ALOG(SF2)VAL2 = B+(D*X2)+(E*(X2**2))+(F*(X2**3))+(G*(X2**4))KV2 = EXP(VAL2)C C CALCULATE THE VAPOR VELOCITY FACTOR FOR HORIZONTAL SEPARATOR KH = 1.25 * KV2VMAX2 = KH*SQRT((DENL2-DENV2)/DENV2) CALCULATE THE LIQUID VOLUMETRIC RATE, ft^3/sec. С QL2 = WL2/(DENL2*3600.0)CALCULATE THE VAPOR VOLUMETRIC RATE, ft^3/sec. С QV2 = WV2/(DENV2 * 3600.0)CALCULATE THE VAPOR FLOW AREA, ft^2. С AV2 = QV2/VMAX2CALCULATE THE TOTAL CROSS-SECTIONAL AREA с OF HORIZONTAL VESSEL. ft^2. С AT = AV2/0.2CALCULATE THE DIAMETER, ft. С D2 = SQRT((4.0*AT)/PI)ASSUMING THE LENGTH TO DIAMETER RATIO = 3 с THE LENGTH TO DIAMETER RATIO, (i.e. RATIO=L/D=3) С $I_{.} = RATIO*D2$ THE SEPARATOR VAPOR AREA IS ASSUMED TO OCCUPY 20% OF THE TOTAL С CROSS-SECTIONAL AREA. THEREFORE THE LIQUID AREA WILL OCCUPY 80% OF С č THE TOTAL CROSS-SECTIONAL AREA. CALCULATE THE FILL LIQUID VOLUME, ft^3. С AL = 0.8 * ATс CALCULATE THE VESSEL VOLUME, ft^3. VESVOL = AT*LCALCULATE THE LIQUID SURGE TIME, min. С CALT2 = (AL*L*DENL2*60.)/WL2 RETURN END C C THIS PROGRAM PRINTS THE RESULTS OF THE VERTICAL SEPARATOR С c SUBROUTINE OUTV REAL KV1 COMMON/DATA1/ WL1, WV1, DENL1, DENV1, T1 COMMON/DATA2/ SF1, KV1, VMAX1, D1 COMMON/DATA3/ UMAX, QL1, QV1, UMIN, V, HL WRITE (1, 100) FORMAT (///,25X, 'VERTICAL SEPARATOR SIZING',/,78(1H*)) 100

110	1 2 3 4 5 6	<pre>WRITE (1, 110) WL1, WV1, DENL1, DENV1, QL1, QV1, T1 FORMAT (5X,'LIQUID FLOWRATE, 1b/hr.:', T60, F10.0,/, 5X,'VAPOR FLOWRATE, 1b/hr.:', T60, F10.0,/, 5X,'VAPOR FLOWRATE, 1b/hr.:', T60, F9.3,/, 5X,'VAPOR DENSITY, 1b/ft^3.:', T60, F9.3,/, 5X,'LIQUID VOLUMETRIC RATE, ft^3/sec.:', T60, F9.4,/, 5X,'VAPOR VOLUMETRIC RATE, ft^3/sec.:', T60, F9.4,/, 5X,'LIQUID SURGE TIME, min.:', T60, F6.1)</pre>
120	1 2 3	WRITE (1, 120) SF1, KV1, VMAX1, D1 FORMAT (5X,'SEPARATION FACTOR:', T60, F8.4,/, 5X,'VAPOR VELOCITY FACTOR:', T60, F8.3,/, 5X,'MAXIMUM VAPOR VELOCITY, ft/s.:',T60, F8.3,/, 5X,'VESSEL DIAMETER, ft:', T60,F8.3)
130	1 2 3	WRITE (1, 130) UMIN, UMAX, V, HL FORMAT (5X,'MINIMUM VAPOR-LIQUID NOZZLE VELOCITY, ft/s.:', T60, F8.3,/,5X,'MAXIMUM VAPOR-LIQUID NOZZLE VELOCITY, ft/s.:', T60, F8.3,/,5X,'REQUIRED VESSEL VOLUME, ft^3:', T60,F8.3,/, 5X,'LIQUID HEIGHT, ft.:', T60, F8.3)
140		WRITE (1, 140) FORMAT (78(1H-))
		RETURN END
c c c		**************************************
		SUBROUTINE OUTH REAL L, KH
		COMMON/DATA4/ WL2, WV2, DENL2, DENV2, RATIO COMMON/DATA5/ SF2, KH, QL2, QV2, VMAX2, D2, L, VESVOL, T2, CALT2
		WRITE(1, 150)
150		<pre>FORMAT(//,25X,'HORIZONTAL SEPARATOR SIZING',/78(1H*))</pre>
160	1 2 3 4 5 6 7	<pre>WRITE (1, 160) WL2, WV2, DENL2, DENV2, QL2, QV2, T2, RATIO FORMAT (5X,'LIQUID FLOWRATE, lb/hr.:', T60, F10.0,/, 5X,'VAPOR FLOWRATE, lb/hr.:', T60, F10.0,/, 5X,'LIQUID DENSITY, lb/ft^3.:', T60, F9.3,/, 5X,'VAPOR DENSITY, lb/ft^3.:', T60, F9.3,/, 5X,'LIQUID VOLUMETRIC RATE, ft^3/sec.:', T60, F9.4,/, 5X,'LIQUID SURGE TIME, min.:', T60, F9.3,/, 5X,'LIQUID SURGE TIME, min.:', T60, F9.3,/, 5X,'LIQUID SURGE TIME, min.:', T60, F9.3,/, 5X,'LENGTH TO DIAMETER RATIO (L/D):', T60, F6.1)</pre>
170	1 2 3	WRITE (1, 170) SF2, KH, QV2, VMAX2 FORMAT (5X,'SEPARATION FACTOR:',T60,F8.3,/, 5X,'VAPOR VELOCITY FACTOR:',T60,F8.3,/, 5X,'VAPOR VOLUMETRIC RATE, ft^3/s.:', T60, F9.3,/, 5X,'MAXIMUM VAPOR VELOCITY, ft/s.:',T60, F8.3)
180	1 2 3	<pre>WRITE (1, 180) D2, L, VESVOL, CALT2 FORMAT (5X,'VESSEL DIAMETER, ft.:',T60, F8.3,/, 5X,'VESSEL LENGTH, ft.:', T60, F8.3,/, 5X,'VESSEL VOLUME, ft^3.:', T60, F9.3,/, 5X,'CALCULATED LIQUID SURGE TIME, min.:', T60, F8.3)</pre>
190		WRITE (1, 190) Format (78(1H-))
с		FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.
		WRITE (1, *) CHAR(12)
		RETURN END

PROGRAM PROG42

С	***************************************
С	THIS PROGRAM CALCULATES THE SEGMENTAL AREA OF A CIRLCE COUPLED
с	WITH A MODIFIED DOOLITTLE FORMULA FOR DISHED HEADS TO CALCULATE
č	PARTIAL VOLUMES OF HORIZONTAL VESSELS.
č	THE PROGRAM WILL CALCULATE PARTIAL VOLUMES FOR HORIZONTAL
с	CYLINDRICAL VESSELS WITH FLAT, DISHED, ELLIPTICAL AND HEMISPHERICAL
с	HEADS, AND VERTICAL CYLINDRICAL TANKS.
С	*****
с	CL = CHORD LENGTH OF SEGMENTAL AREA, in.
č	$AC = AREA OF CHORD, in^2.$
č	$VHC = VOLUME OF HORIZONTAL CYLINDER, in^3.$
с	VDH = VOLUME OF DISHED HEADS, in^3.
С	VELL = VOLUME OF ELLIPTICAL HEADS, in^3.
С	VHS = VOLUME OF HEMISPHERICAL HEADS, in^3.
с	<pre>VT = TOTAL VOLUME OF HORIZONTAL VESSEL, in^3.</pre>
c	D = VESSEL, TANK, OR CIRCLE DIAMETER, in.
č	H = CHORD OR LIQUID HEIGHT, in.
с	R = RADIUS OF VESSEL OR TANK, in.
с	L = VESSEL LENGTH, in.
с	VH = HEAD VOLUMES, in^3.
с	BBL = VOLUME IN BARRELS
с	$AREA = CIRCLE AREA, in^2.$
č	*****
C	
	REAL L1, L2, L3, L4, L5
	COMMON PI
	COMMON/DATA1/ D1, H1, L1
	COMMON/DATA2/ D2, H2, L2
	COMMON/DATA3/ D3, H3, L3
	COMMON/DATA4/ D4, H4, L4
	COMMON/DATA5/ D5, H5, L5
	COMMON/PROG1/ VHC, VHCCFT, VHCGAL, VHCBBL
	COMMON/PROG2/ VT, VTCFT, VTGAL, VTBBL
	COMMON/PROG3/ VTELL, VTECFT, VTEGAL, VTEBBL
	COMMON/PROG4/ VTHS, VTHCFT, VTHGAL, VTHBBL
	COMMON/PROG5/ VVT, VVCFT, VVGAL, VVBBL
	COMMON/FROGS/ VVI, VVCFI, VVGRL, VVBBL
	OPEN (UNIT=3, FILE='DATA42.DAT', STATUS='OLD', ERR=18)
	OPEN (UNIT=1, FILE='PRN')
	PI = 3.1415927
5	PEAD (2 + FDD-10) T
5	READ (3, *, ERR=19) I
	GO TO (11, 22, 33, 44, 55), I
с	READ TANK DIAMETER, D1: in.
c	READ LIQUID DEPTH, H1: in.
č	READ TANK LENGTH, L1: in.
~	
11	DEAD (3 + E0D-10) UI II
11	READ (3, *, ERR=19) D1, H1, L1
	GO TO 66

GO TO 66

C C C	READ TANK DIAMETER, D2: in. READ LIQUID DEPTH, H2: in. READ TANK LENGTH, L2: in.
22	READ (3, *, ERR=19) D2, H2, L2
	GO TO 77
C C C	READ TANK DIAMETER, D3: in. READ LIQUID DEPTH, H3: in. READ TANK LENGTH, L3: in.
33	READ (3, *, ERR=19) D3, H3, L3
	GO TO 88
C C C	READ TANK DIAMETER, D4: in. READ LIQUID DEPTH, H4: in. READ TANK LENGTH, L4: in.
44	READ (3, *, ERR=19) D4, H4, L4
	GO TO 99
55	READ (3, *, ERR=19) D5, H5, L5
	GO TO 111
18 12	WRITE (*, 12) Format (6X, 'DATA FILE DOES NOT EXIST') GO TO 999
19 13	WRITE (*, 13) Format(6X, 'Error Message in the data value') Go to 999
с	CALCULATE THE VOLUME OF THE TANK WITH FLAT HEADS
66	CALL FLAT
с	PRINT THE RESULTS OF THE TANK WITH FLAT HEADS
	CALL OFLAT
	GO TO 5
с	CALCULATE THE VOLUME OF THE TANK WITH DISHED HEADS
77	CALL DISHED
с	PRINT THE RESULTS OF THE TANK WITH DISHED HEADS
	CALL ODISH
	GO TO 5
с	CALCULATE THE VOLUME OF THE TANK WITH ELLIPTICAL HEADS
00	CALL FILTD

88 CALL ELLIP

с PRINT THE RESULTS OF THE TANK WITH ELLIPTICAL HEADS. CALL OELLIP GO TO 5 CALCULATE THE VOLUME OF THE TANK WITH HEMISPHERICAL HEADS С 99 CALL HEMISP с PRINT THE RESULTS OF THE TANK WITH HEMISPHERICAL HEADS. CALL OHEMIS GO TO 5 CALCULATE THE PARTIAL VOLUME OF A VERTICAL TANK С 111 CALL VERT PRINT THE RESULTS OF THE PARTIAL VOLUME OF A VERTICAL С С VESSEL CALL OVERT GO TO 999 CLOSE (3, STATUS='KEEP') CLOSE (1) 999 STOP END 0000 ********************************* THIS PROGRAM CALCULATES THE PARTIAL VOLUME FOR A HORIZONTAL CYLINDRICAL VESSEL WITH FLAT HEADS SUBROUTINE FLAT REAL L1 COMMON PI COMMON/DATA1/ D1, H1, L1 COMMON/PROG1/ VHC, VHCCFT, VHCGAL, VHCBBL R1 = D1/2. CL1 = 2.*SQRT(D1*H1-(H1**2)) CLI = 2.*SQRT(DI*n1=(h1**2)) VAL1 = (R1-H1)/R1 VAL2 = (R1-H1)*SQRT(2.*R1*H1-(H1**2)) AC1 = ((R1**2)*ACOS(VAL1))-VAL2 AREA1 = (0.25*PI*D1**2) VHC = AC1*L1 С CONVERT THE RESULTS TO FT^3, GALLONS AND BARRELS CALL CONVT (VHC, VHCCFT, VHCGAL, VHCBBL)

RETURN

END

```
0000
                     THIS PROGRAM CALCULATES THE PARTIAL VOLUME FOR A HORIZONTAL
                     CYLINDRICAL VESSEL WITH DISHED HEADS
                      *****************
                     SUBROUTINE DISHED
                     REAL L2
                     COMMON PT
                     COMMON/DATA2/ D2, H2, L2
COMMON/PROG2/ VT, VTCFT, VTGAL, VTBBL
                     R2 = D2/2.
                     CL2 = 2.*SQRT(D2*H2-(H2**2))
                     \begin{aligned} & CL2 &= 2 \cdot SQR1(12^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L^{-1}L
                      VHC2 = AC2 * L2
                     VDH = 0.215483*H2*(1.5*D2-H2)
                     VT = 2.*VDH+VHC2
                     CONVERT THE RESULTS TO CUBIC FEET, GALLONS AND BARRELS.
с
                     CALL CONVT (VT, VTCFT, VTGAL, VTBBL)
                     RETURN
                      END
                      CCCC
                      THIS PROGRAM CALCULATES THE PARTIAL VOLUME FOR A HORIZONTAL
                     CYLINDRICAL VESSEL WITH ELLIPTICAL HEADS
                                                                                                                               ....
                     SUBROUTINE ELLIP
                     REAL L3
                      COMMON PI
                      COMMON/DATA3/ D3, H3, L3
COMMON/PROG3/VTELL,VTECFT,VTEGAL,VTEBBL
                     R3 = D3/2.
CL3 = 2.*SQRT(D3*H3-(H3**2))
                      VAL5 = (R3-H3)/R3
VAL6 = (R3-H3)*SQRT(2.*R3*H3~(H3**2))
                      AC3 = ((R3**2)*ACOS(VAL5))-VAL6
                      VHC3 = AC3 * L3
                      VELL = 0.5236*(H3**2)*(1.5*D3-H3)
                      VTELL = 2.*VELL+VHC3
¢
                      CONVERT THE RESULTS TO CUBIC FEET, GALLONS AND BARRELS.
                      CALL CONVT (VTELL, VTECFT, VTEGAL, VTEBBL)
                     RETURN
                     END
с
```

```
с
с
с
       THIS PROGRAM CALCULATES THE PARTIAL VOLUME FOR A HORIZONTAL
       CYLINDRICAL VESSEL WITH HEMISPHERICAL HEADS
       SUBROUTINE HEMISP
       REAL L4
       COMMON PI
       COMMON/DATA4/ D4, H4, L4
COMMON/PROG4/ VTHS, VTHCFT, VTHGAL, VTHBBL
       R4 = D4/2.
       CL4 = 2.*SQRT(D4*H4-(H4**2))
       VAL7 = (R4-H4)/R4

VAL8 = (R4-H4)*SQRT(2.*R4*H4-(H4**2))
       AC4 = ((R4**2)*ACOS(VAL7))-VAL8
VHC4 = AC4*L4
       VELL4 = 0.5236*(H4**2)*(1.5*D4-H4)
VHS = 2.*VELL4
VTHS = 2.*VHS+VHC4
с
       CONVERT THE RESULTS TO CUBIC FEET, GALLONS AND BARRELS.
       CALL CONVT (VTHS, VTHCFT, VTHGAL, VTHBBL)
       RETURN
       END
С
       ******************
                                 .................
C
C
       THIS PROGRAM CALCULATES THE PARTIAL VOLUME FOR A VERTICAL
       CYLINDRICAL VESSEL
с
       *******
       SUBROUTINE VERT
       COMMON PI
       COMMON/DATA5/ D5, H5, L5
COMMON/PROG5/ VVT, VVCFT, VVGAL, VVBBL
       VVT = 0.25*PI*(D5**2)*H5
С
       CONVERT THE RESULTS TO CUBIC FEET, GALLONS AND BARRELS.
       CALL CONVT (VVT, VVCFT, VVGAL, VVBBL)
       RETURN
       END
с
                 C
C
C
C
C
       THIS PROGRAM CONVERTS THE RESULTS TO CUBIC FEET, GALLONS, AND
       BARRELS.
       *****
       SUBROUTINE CONVT (VOL, VLFT3, VLGAL, VLBAR)
       VLFT3 = VOL/1728.0
```

310

```
VLGAL = VOL*0.004329
         VLBAR = VOL*0.00010307
         RETURN
         END
С
         THIS PROGRAM PRINTS THE RESULTS OF THE TANK VOLUME WITH FLAT
C
C
C
         HEADS
         SUBROUTINE OFLAT
         REAL L1
         COMMON/DATA1/ D1, H1, L1
COMMON/PROG1/ VHC, VHCCFT, VHCGAL, VHCBBL
         WRITE (1, 100)
         FORMAT (//,25X, 'TANK VOLUME WITH FLAT HEADS',/,78(1H*))
 100
         WRITE (1, 110) D1, H1, L1
         FORMAT (5X,'TANK DIAMETER, in.:', T60, F9.3,/,
5X,'LIQUID DEPTH, in.:', T60, F9.3,/,
5X,'TANK LENGTH, in.:', T60, F9.3)
 110
     1
     2
        WRITE (1, 120) VHC, VHCCFT, VHCGAL, VHCBBL
FORMAT (5x,'TANK VOLUME, in.^3:', T60, E14.6,/,
5x,'TANK VOLUME, ft.^3:', T60, E14.6,/,
5x,'TANK VOLUME, gal:', T60, E14.6,/,
5x,'TANK VOLUME, bbl:', T60, E14.6)
 120
     1
     2
     3
         WRITE (1, 130)
FORMAT (78(1H-))
 130
         RETURN
         END
         0000
         THIS PROGRAM PRINTS THE RESULTS OF THE TANK VOLUME WITH DISHED
         HEADS.
         SUBROUTINE ODISH
         REAL L2
         COMMON/DATA2/ D2, H2, L2
COMMON/PROG2/ VT, VTCFT, VTGAL, VTBBL
         WRITE (1, 140)
 140
         FORMAT (//,25X,'TANK VOLUME WITH DISHED HEADS',/,78(1H*))
         WRITE (1, 150) D2, H2, L2
FORMAT (5X, TANK DIAMETER, in.:', T60, F9.3,/,
5X,'LIQUID DEPTH, in.:', T60, F9.3,/,
5X,'TANK LENGTH, in.:', T60, F9.3)
 150
     1
      2
```

```
WRITE (1, 160) VT, VTCFT, VTGAL, VTBBL
FORMAT (5x,'TANK VOLUME, in^3:', T60, E14.6,/,
5x,'TANK VOLUME, ft.^3:', T60, E14.6,/,
5x,'TANK VOLUME, gal:', T60, E14.6,
5x,'TANK VOLUME, bbl:', T60, E14.6)
 160
      1
      2
      3
          WRITE (1, 170)
FORMAT (78(1H-))
 170
          RETURN
          END
          ***************
с
                                                                                **********
c
c
c
          THIS PROGRAM PRINTS THE RESULTS OF THE TANK VOLUME WITH
          ELLIPTICAL HEADS
          ************
                                    SUBROUTINE OELLIP
          REAL L3
          COMMON/DATA3/ D3, H3, L3
COMMON/PROG3/ VTELL, VTECFT, VTEGAL, VTEBBL
          WRITE (1, 180)
 180
          FORMAT (//,25X, 'TANK VOLUME WITH ELLIPTICAL HEADS',/,78(1H*))
          WRITE (1, 190) D3, H3, L3
          FORMAT (5X, 'TANK DIAMETER, in.:', T60, F9.3,/,
5X,'LIQUID DEPTH, in.:', T60, F9.3,/,
5X,'TANK LENGTH, in.:', T60, F9.3)
 190
      1
      2
          WRITE (1, 200) VTELL, VTECFT, VTEGAL, VTEBBL
FORMAT (5X,'TANK VOLUME, in.^3:', T60, E14.6,/,
5X,'TANK VOLUME, ft.^3:', T60, E14.6,/,
5X,'TANK VOLUME, gal:', T60, E14.6,/
5X,'TANK VOLUME, bbl:', T60, E14.6)
 200
      1
      2
      3
          WRITE (1, 210)
          FORMAT (78(1H-))
 210
          RETURN
          END
с
          *************************
                                                                            *************
С
          THIS PROGRAM PRINTS THE RESULTS OF THE TANK VOLUME FOR
С
          HEMISPHERICAL HEADS.
с
          SUBROUTINE OHEMIS
          REAL L4
          COMMON/DATA4/ D4, H4, L4
COMMON/PROG4/ VTHS, VTHCFT, VTHGAL, VTHBBL
          WRITE (1, 220)
          FORMAT (//,25X,'TANK VOLUME WITH HEMISPHERICAL HEADS',/,78(1H*))
 220
```

230 1 2	WRITE (1, 230) D4, H4, L4 FORMAT (5X,'TANK DIAMETER, in.:', T60, F9.3,/, 5X,'LIQUID DEPTH, in.:', T60, F9.3,/, 5X,'TANK LENGTH, in.:', T60, F9.3)
240 1 2 3	<pre>WRITE (1, 240) VTHS, VTHCFT, VTHGAL, VTHBBL FORMAT (5X,'TANK VOLUME, in.^3:', T60, E14.6,/, 5X,'TANK VOLUME, ft.^3:', T60, E14.6,/, 5X,'TANK VOLUME, gal:', T60, E14.6,/, 5X,'TANK VOLUME, bbl:', T60, E14.6)</pre>
250	WRITE (1, 250) Format (78(1H-))
	RETURN END
0000	THIS PROGRAM PRINTS THE RESULTS OF THE PARTIAL VOLUME FOR A VERTICAL CYLINDRICAL VESSEL
	SUBROUTINE OVERT
	REAL L5 COMMON/DATA5/ D5, H5, L5 COMMON/PROG5/ VVT, VVCFT, VVGAL, VVBBL
260	WRITE (1, 260) FORMAT (//,25X,'VERTICAL TANK VOLUME', /, 78(1H*))
270 1 2	WRITE (1, 270) D5, H5, L5 FORMAT (5X,'TANK DIAMETER, in.:', T60, F9.3,/, 5X,'LIQUID DEPTH, in.:', T60, F9.3,/, 5X,'TANK LENGTH, in.:', T60, F9.3)
280 1 2 3	WRITE (1, 280) VVT, VVCFT, VVGAL, VVBBL FORMAT (5X,'TANK VOLUME, in.^3:', T60, E14.6,/, 5X,'TANK VOLUME, ft.3:', T60, E14.6,/, 5X,'TANK VOLUME, gal:', T60, E14.6,/, 5X,'TANK VOLUME, bbl:', T60, E14.6)
290	WRITE (1, 290) Format (78(1H-))
с	FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR(12)
	RETURN END

PROGRAM PROG43

с	************
č	A PROGRAM FOR PRELIMINARY VESSEL DESIGN.
č	THIS PROGRAM WILL ESTIMATE SHELL THICKNESS AND WEIGHT,
c	HEAD THICKNESS AND WEIGHT FOR CYLINDRICAL PRESSURE VESSELS.
	IN ADDITION, HEAD THICKNESS AND WEIGHT FOR CHEINDRICHE FRESSORE VESSERS.
c	
с	HEMISPHERICAL, ELLIPSOIDAL AND TORISPHERICAL HEADS. TOTAL VESSEL
с	WEIGHT IS CALCULATED (SHELL AND HEADS) EXCLUSIVE OF NOZZLES,
с	ATTACHMENTS AND VESSEL INTERNALS.
с	***************************************
с	TKS = SHELL THICKNESS, in.
С	TKHH = HEMISPHERICAL HEAD THICKNESS, in.
с	TKELL = ELLIPSOIDAL HEAD THICKNESS, in.
с	TKTOR = TORISPHERICAL HEAD THICKNESS, in.
с	TKELL = ELLIPSOIDAL HEAD THICKNESS, in. TKTOR = TORISPHERICAL HEAD THICKNESS, in. P = INTERNAL DESIGN PRESSURE, psig. R = INSIDE RADIUS, in.
с	R = INSIDE RADIUS, in.
с	$S = ALLOWABLE STRESS, 1b/in^2.$
с	S = ALLOWABLE STRESS, lb/in^2. E = JOINT EFFICIENCY
с	C = CORROSION ALLOWANCE, in.
с	ID = INSIDE DIAMETER OF SHELL OR HEAD, in. RC = CROWN RADIUS OF TORISPHERICAL HEAD, in.
с	RC = CROWN RADIUS OF TORISPHERICAL HEAD, in.
с	LN = SHELL LENGTH, ft.
с	DENV = DENSITY OF VESSEL MATERIAL, lb/ft^3. OD = OUTSIDE DIAMETER OF HEAD, in. HFAC = HEAD BLANK DIAMETER FACTOR SF = STRAIGHT FLANGE LENGTH, in. BD = HEAD BLANK DIAMETER, in.
с	OD = OUTSIDE DIAMETER OF HEAD, in.
с	HFAC = HEAD BLANK DIAMETER FACTOR
c	SF = STRAIGHT FLANGE LENGTH, in.
c	SF = STRAIGHT FLANGE LENGTH, in. BD = HEAD BLANK DIAMETER, in.
č	WTH = WEIGHT OF HEAD, 1b
c	WTS = WEIGHT OF VESSEL SHELL, 1b.
c	TOTWT = TOTAL VESSEL WEIGHT, 1b.
ċ	*******
	REAL LN1, LN2, LN3, ID1, ID2, ID3
	COMMON PI
	COMMON/DATA1/ ID1, LN1, P1, SF1, C1, DENS1, ES1, EH1, S1
	COMMON/RES1/ TKS1, TKH1, WTS1, WTH1, TOTWT1
	COMMON/DATA2/ ID2, LN2, P2, SF2, C2, DENS2, ES2, EH2, S2
	COMMON/RES2/ TKS2, TKHS2, WTS2, WTH2, TOTWT2
	COMMON/DATA3/ ID3, LN3, P3, SF3, C3, DENS3, ES3, EH3, S3, RC
	COMMON/RES3/ TKS3, TKHS3, WTS3, WTH3, TOTWT3
	OPEN (UNIT=3, FILE='DATA43.DAT', STATUS='OLD', ERR=18)
	OPEN (UNIT=1, FILE='PRN')
	PI = 3.1415927
5	READ (3, *, ERR=19) I
	GO TO (11, 22, 33), I
с	READ INTERNAL DIAMETER, ID1: in.
с	READ SHELL LENGTH, LN1: in.
с	READ INTERNAL DESIGN PRESSURE, P1: psig.
~	DEAD CODATCHO FLANCE LENCON CEL.

C READ STRAIGHT FLANGE LENGTH, SF1: in.

0 0 0 0 0 0 0 0	READ CORROSION ALLOWANCE, C1: in. READ DENSITY OF VESSEL MATERIAL, DENS1: lb/ft^3. READ JOINT EFFICIENCY OF SHELL, ES1 READ JOINT EFFICIENCY OF HEAD, EH1 READ ALLOWABLE STRESS, S1: lb/in^2.
11	READ (3, *, ERR=19) ID1, LN1, P1 READ (3, *, ERR=19) SF1, C1, DENS1 READ (3, *, ERR=19) ES1, EH1, S1
	GO TO 44
0 0 0 0 0 0 0 0 0	READ INTERNAL DIAMETER, ID2: in. READ SHELL LENGTH, LN2: in. READ INTERNAL DESIGN PRESSURE, P2: psig. READ STRAIGHT FLANGE LENGTH, SF2: in. READ CORROSION ALLOWANCE, C2: in. READ DENSITY OF VESSEL MATERIAL, DENS2: lb/ft^3. READ JOINT EFFICIENCY OF SHELL, ES2: READ JOINT EFFICIENCY OF HEAD, EH2: READ ALLOWABLE STRESS, S2: lb/in^2.
22	READ (3, *, ERR=19) ID2, LN2, P2 READ (3, *, ERR=19) SF2, C2, DENS2 READ (3, *, ERR=19) ES2, EH2, S2
	GO TO 55
с с с с с с с с	READ INTERNAL DIAMETER, ID3: in. READ SHELL LENGTH, LN3: in. READ INTERNAL DESIGN PRESSURE, P3: psig. READ STRAIGHT FLANCE LENGTH, SF3: in. READ CORROSION ALLOWANCE, C3: in. READ DENSITY OF VESSEL MATERIAL, DENS3: lb/ft^3. READ JOINT EFFICIENCY OF SHELL, ES3: READ JOINT EFFICIENCY OF HEAD, EH3: READ CROWN RADIUS OF TORISPHERICAL HEAD, RC: in.
33	READ (3, *, ERR=19) ID3, LN3, P3 READ (3, *, ERR=19) SF3, C3, DENS3 READ (3, *, ERR=19) ES3, EH3, S3, RC
	GO TO 66
18 13	WRITE (*, 13) Format (6x, 'data file does not exist') Go to 999
19 14	WRITE (*, 14) Format (6x, 'error message in the data value') Go to 999
c c	CALCULATE THE SHELL THICKNESS, WEIGHT AND HEAD THICKNESS FOR HEMISPHERICAL HEADS.
44	CALL HEMISP

с PRINT THE RESULTS OF THE VESSEL WITH HEMISPHERICAL HEADS. CALL OHEMISP GO TO 5 CALCULATE THE SHELL THICKNESS, WEIGHT AND HEAD THICKNESS FOR c c ELLIPSOIDAL HEADS. 55 CALL ELLIP с PRINT THE RESULTS OF THE VESSEL WITH ELLIPSOIDAL HEADS. CALL OELLIP GO TO 5 CALCULATE THE SHELL THICKNESS, WEIGHT AND HEAD THICKNESS FOR С С TORISPHERICAL HEADS. CALL TORIS 66 PRINT THE RESULTS OF THE VESSEL WITH TORISPHERICAL HEADS. с CALL OTORIS GO TO 999 CLOSE (3, STATUS='KEEP') CLOSE (1) 999 STOP END с с с с VESSEL DESIGN WITH HEMISPHERICAL HEADS ***** SUBROUTINE HEMISP REAL LN1, ID1, OD1 COMMON PI COMMON/DATA1/ ID1, LN1, P1, SF1, C1, DENS1, ES1, EH1, S1 COMMON/RES1/ TKS1, TKH1, WTS1, WTH1, TOTWT1 с CALCULATE SHELL THICKNESS, in R1 = ID1/2. TKS1 = (P1*R1)/(S1*ES1-0.6*P1)+C1 OD1 = ID1 + 2.0*TKS1 с CALCULATE WEIGHT OF VESSEL SHELL, 1b WTS1 = ((OD1+TKS1)*PI*LN1*TKS1*DENS1)/144. С CALCULATE WEIGHT OF HEAD, 1b

TKH1 = (P1*R1)/(2.*S1*EH1-0.2*P1)

```
RAT1 = OD1/TKH1
       IF ((OD1/TKH1) .GE. 10.0 .AND. (OD1/TKH1) .LT. 18.0) THEM
       HFAC = 1.70
       ELSEIF ((OD1/TKH1) .GE. 18.0 .AND. (OD1/TKH1) .LT. 30.0) THEN
       HFAC = 1.65
       ELSEIF ((OD1/TKH1) .GT. 30.0 ) THEN
       HFAC = 1.60
       ENDIF
       BD1 = (OD1*HFAC+2.*SF1)/12.
       WTH1 = (0.25*(BD1**2)*PI*TKH1*DENS1*2.)/12.
С
       CALCULATE TOTAL VESSEL WEIGHT
       TOTWT1 = WTS1 + WTH1
       RETURN
       END
С
       VESSEL DESIGN WITH ELLIPSOIDAL HEADS
с
ċ
       SUBROUTINE ELLIP
       REAL LN2, ID2, OD2
       COMMON PI
       COMMON/DATA2/ ID2, LN2, P2, SF2, C2, DENS2, ES2, EH2, S2
COMMON/RES2/ TKS2, TKH2, WTS2, WTH2, TOTWT2
с
       CALCULATE SHELL THICKNESS, in
       R2 = ID2/2.
       TKS2 = (P2*R2)/(S2*ES2-0.6*P2)+C2
       OD2 = ID2 + 2.0 * TKS2
       CALCULATE WEIGHT OF VESSEL SHELL, 1b
С
       WTS2 = ((OD2+TKS2)*PI*LN2*TKS2*DENS2)/144.
С
       CALCULATE WEIGHT OF HEAD, 1b
       TKH2 = (P2*ID2)/(2.*S2*EH2-0.2*P2)+C2
       RAT2 = OD2/TKH2
       IF ((OD2/TKH2) .GE. 10.0 .AND. (OD2/TKH2) .LT. 20.0) THEN
       HFAC = 1.30
       ELSEIF ((OD2/TKH2) .GT. 20.0) THEN
       HFAC = 1.24
       ENDIF
       BD2 = (OD2*HFAC+2.*SF2)/12.
       WTH2 = (0.25*(BD2**2)*PI*TKH2*DENS2*2.)/12.0
```

C CALCULATE TOTAL WEIGHT

```
TOTWT2 = WTS2 + WTH2
       RETURN
       END
       ¢
С
                     VESSEL DESIGN WITH TORISPHERICAL HEADS
¢
       SUBROUTINE TORIS
       REAL LN3, ID3, OD3
       COMMON PI
       COMMON/DATA3/ ID3, LN3, P3, SF3, C3, DENS3, ES3, EH3, S3, RC
COMMON/RES3/ TKS3, TKH3, WTS3, WTH3, TOTWT3
с
       CALCULATE SHELL THICKNESS, in
       R3 = ID3/2.
       TKS3 = (P3*R3)/(S3*ES3-0.6*P3)+C3
       OD3 = ID3 + 2.0 * TKS3
       CALCULATE WEIGHT OF VESSEL SHELL, 1b
с
       WTS3 = ((OD3+TKS3)*PI*LN3*TKS3*DENS3)/144.
C
       CALCULATE WEIGHT OF HEAD, 1b
       TKH3 = ((0.885*P3*RC)/(S3*EH3-0.1*P3))+C3
       RAT3 = OD3/TKH3
       IF ((OD3/TKH3) .GE. 20.0 .AND. (OD3/TKH3) .LT. 30.0) THEN
       HFAC = 1.15
       ELSEIF ((OD3/TKH3) .GE. 30.0 .AND. (OD3/TKH3) .LT. 50.0) THEN
       HFAC = 1.11
       ELSEIF ((OD3/TKH3) .GT. 50.0) THEN
       HFAC = 1.09
       ENDIF
       BD3 = (OD3 * HFAC + 2.0 * SF3)/12.
       WTH3 = (0.25*(BD3**2)*PI*TKH3*DENS3*2.)/12.
с
       CALCULATE TOTAL WEIGHT
       TOTWT3 = WTS3 + WTH3
       RETURN
       END
CCCC
       ******************
                                   ******
                                                ******
       THIS PROGRAM PRINTS THE RESULTS OF THE VESSEL WITH
       HEMISPHERICAL HEADS.
       SUBROUTINE OHEMISP
```

REAL ID1, LN1

	COMMON/DATA1/ ID1, LN1, P1, SF1, C1, DENS1, ES1, EH1, S1 COMMON/RES1/ TKS1, TKH1, WTS1, WTH1, TOTWT1
100 *	WRITE (1, 100) FORMAT (//,25X, 'VESSEL DESIGN WITH HEMISPHERICAL HEADS', /,78(1H*))
110 1 2	WRITE (1, 110) ID1, LN1, P1 FORMAT (5X,'INTERNAL DIAMETER, in.:', T60, F9.3,/, 5X,'SHELL LENGHT, ft.:', T60, F9.3,/, 5X,'INTERNAL DESIGN PRESSURE, psig:', T60, F9.3)
120 1 2	<pre>WRITE (1, 120) SF1, C1, DENS1 FORMAT (5X,'STRAIGHT FLANGE LENGTH, in.:', T60, F9.3,/, 5X,'CORROSION ALLOWANCE, in.:', T60, F9.3,/, 5X,'DENSITY OF VESSEL MATERIAL, lb/ft^3.:', T60, F9.3)</pre>
130 1 2	<pre>WRITE (1, 130) ES1, EH1, S1 FORMAT (5X,'JOINT EFFICIENCY OF SHELL:', T60, F9.3,/, 5X,'JOINT EFFICIENCY OF HEAD:', T60, F9.3,/, 5X,'ALLOWABLE STRESS, lb/in^2.:', T60, F9.3)</pre>
140 1 2 3 4	<pre>WRITE (1, 140) TKS1, TKH1, WTS1, WTH1, TOTWT1 FORMAT (5X,'CALCULATED SHELL THICKNESS, in.:', T60, F9.3,/, 5X,'CALCULATED HEAD THICKNESS, in.:', T60, F9.3,/, 5X,'WEIGHT OF VESSEL SHELL, 1b:', T60, F12.2,/, 5X,'WEIGHT OF VESSEL HEAD, 1b:', T60, F12.2,/, 5X,'TOTAL VESSEL WEIGHT, 1b:', T60, F12.2)</pre>
150	WRITE (1, 150) Format (78(1H-))
	RETURN END
с с с с	**************************************
	SUBROUTINE OELLIP
	REAL ID2, LN2 COMMON/DATA2/ID2, LN2, P2, SF2, C2, DENS2, ES2, EH2, S2 COMMON/RES2/TKS2, TKH2, WTS2, WTH2, TOTWT2
160 1	WRITE (1, 160) FORMAT (//,25X, 'VESSEL DESIGN WITH ELLIPSOIDAL HEADS', /,78(1H*))
170 1 2	WRITE (1, 170) ID2, LN2, P2 FORMAT (5X, 'INTERNAL DIAMETER, in.:', T60, F9.3,/, 5X,'SHELL LENGTH, ft.:', T60, F9.3,/, 5X,'INTERNAL DESIGN PRESSURE, psig.:', T60, F9.3)

180	1 2	<pre>WRITE (1, 180) SF2, C2, DENS2 FORMAT (5X,'STRAIGHT FLANGE LENGTH, in.:', T60, F9.3,/, 5X,'CORROSION ALLOWANCE, in.:', T60, F9.3,/, 5X,'DENSITY OF VESSEL MATERIAL, lb/ft^3.:',T60, F9.3)</pre>			
190	1 2	<pre>wRITE (1, 190) ES2, EH2, S2 FORMAT (5X,'JOINT EFFICIENCY OF SHELL:', T60, F9.3,/, 5X,'JOINT EFFICIENCY OF HEAD:', T60, F9.3,/, 5X,'ALLOWABLE STRESS, lb/in^2.:', T60, F9.3)</pre>			
200	1 2 3 4	<pre>WRITE (1, 200) TKS2, TKH2, WTS2, WTH2, TOTWT2 FORMAT (5X,'CALCULATED SHELL THICKNESS, in.:', T60, F9.3,/, 5X,'CALCULATED HEAD THICKNESS, in.:', T60, F9.3,/, 5X,'WEIGHT OF VESSEL SHELL, lb:', T60, F12.2,/, 5X,'WEIGHT OF VESSEL HEAD, lb:', T60, F12.2,/, 5X,'TOTAL VESSEL WEIGHT, lb:', T60, F12.2)</pre>			
210		WRITE (1, 210) Format (78(1H-))			
		RETURN END			
с с с с		**************************************			
		SUBROUTINE OTORIS			
		REAL ID3, LN3 COMMON/DATA3/ ID3, LN3, P3, SF3, C3, DENS3, ES3, EH3, S3, RC COMMON/RES3/ TKS3, TKH3, WTS3, WTH3, TOTWT3			
220	1	WRITE (1, 220) FORMAT (//,25X,'VESSEL DESIGN WITH TORISPHERICAL HEADS', /,78(1H*))			
230	1 2	WRITE (1, 230) ID3, LN3, P3 FORMAT (5X,'INTERNAL DIAMETER, in.:', T60, F9.3,/, 5X,'SHELL LENGTH, ft.:', T60, F9.3,/, 5X,'INTERNAL DESIGN PRESSURE, psig.:', T60, F9.3)			
240	1 2	<pre>WRITE (1, 240) SF3, C3, DENS3 FORMAT (5X,'STRAIGHT FLANGE LENGTH, in.:', T60, F9.3,/, 5X,'CORROSION ALLOWANCE, in.:', T60, F9.3,/, 5X,'DENSITY OF VESSEL MATERIAL, lb/ft^3.:', T60, F9.3)</pre>			
250	1 2	WRITE (1, 250) ES3, EH3, S3 FORMAT (5X,'JOINT EFFICIENCY OF SHELL:', T60, F9.3,/, 5X,'JOINT EFFICIENCY OF HEAD:', T60, F9.3,/, 5X,'ALLOWABLE STREES, lb/in.^2:', T60, F9.3)			
260		WRITE (1, 260) TKS3, TKH3, WTS3, WTH3, TOTWT3 FORMAT (5X,'CALCULATED SHELL THICKNESS, in.:', T60, F9.3,/,			
	1 2 3 4	5X,'CALCULATED HEAD THICKNESS, ft.:', T60, F9.3,/, 5X,'WEIGHT OF VESSEL SHELL, lb:', T60, F12.2,/, 5X,'WEIGHT OF VESSEL HEAD, lb:', T60, F12.2,/, 5X,'TOTAL VESSEL WEIGHT, lb:', T60, F12.2)			
270		WRITE (1, 270) FORMAT (78(1H-))			
С		FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.			
		WRITE (1, *) CHAR(12)			
		RETURN END			

PROGRAM PROG44

0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	THE PROGRAM CALCULATES CYCLONE EFFICIENCY BASED ON CYCLONE DIMENSIONS AND ON FLUID AND PARTICLE CHARACTERISTICS. IN ADDITION, THE PROGRAM CALCULATES THE CYCLONE PRESSURE DROP. ************************************
	REAL L, NH, NT COMMON/RES1/ L, D, NH, VI, VSA, RATIO, G, VE, RT, EFF, DELP COMMON/RES2/ NT, DPCRIT, DPMIC, RATIO1 OPEN (UNIT=3, FILE='DATA44.DAT', STATUS='OLD', ERR=18) OPEN (UNIT=1, FILE='PRN')
000000000000000000000000000000000000000	READ CYCLONE OVERALL HEIGHT, H: ft. READ CYCLONE CYLINDRICAL HEIGHT, H1: ft. READ CYCLONE DIAMETER, DC: ft. READ CYCLONE DUST-OUTLET DIAMETER, B: ft. READ CYCLONE GAS INLET LENGTH, A1: ft. READ CYCLONE GAS INLET WIDTH, B1: ft. READ CYCLONE GAS OUTLET LENGTH, S: ft. READ CYCLONE GAS OUTLET LENGTH, S: ft. READ CYCLONE GAS OUTLET DIAMETER, DE: ft. READ CYCLONE GAS OUTLET DIAMETER, DE: ft. READ TOTAL GAS FLOWRATE, Q: ft^3/sec. READ GAS DENSITY, DENF: lb/ft^3. READ GAS DENSITY, VIS: lb/ft.sec. READ GAS TEMPERATURE, T: oF. READ PARTICLE SIZE EQUIVALENT, DPI: ft.
18 20	READ (3, *, ERR=19) H, H1, DC, B READ (3, *, ERR=19) A1, B1, S, DE READ (3, *, ERR=19) Q, DENP, DENF, VIS READ (3, *, ERR=19) T, DPI GO TO 10 WRITE (*, 20) FORMAT (6X, 'DATA FILE DOES NOT EXIST')

```
GO TO 999
       WRITE (*, 25)
FORMAT (6X, 'ERROR MESSAGE IN THE DATA VALUE')
 19
 25
       GO TO 999
С
       CALCULATE THE CYCLONE PRESSURE DROP AND EFFICIENCY
       CALL CALC (H,H1,DC,B,A1,B1,S,DE,Q,DENP,DENF,VIS,T,DPI)
10
       PRINT THE RESULTS OF THE CYCLONE DESIGN.
С
       CALL OUTPUT (H,H1,DC,B,A1,B1,S,DE,Q,DENP,DENF,VIS,T,DPI)
       GO TO 999
       CLOSE (3, STATUS='KEEP')
       CLOSE (1)
999
       STOP
       END
C
C
C
       THIS PROGRAM CALCULATES THE CYCLONE PRESSURE DROP AND EFFICIENCY
       SUBROUTINE CALC(H,H1,DC,B,A1,B1,S,DE,Q,DENP,DENF,VIS,T,DPI)
       PARAMETER (PI=3.1415927, GC=32.2)
       REAL L, NH, NT
       COMMON/RES1/ L, D, NH, VI, VSA, RATIO, G, VE, RT, EFF, DELP
COMMON/RES2/ NT, DPCRIT, DPMIC, RATIO1
с
       CALCULATE L- NATURAL LENGTH i.e. THE DISTANCE BELOW THE GAS
С
       OUTLET WHERE THE VORTEX TURNS, ft.
       L = 2.3*DE*(((DC**2)/(A1*B1))**(1.0/3.0))
       CALCULATE D-DIAMETER OF CENTRAL CORE AT POINT WHERE VORTEX TURNS
С
       D = DC-((DC-B)*((S+L-H1)/(H-H1)))
с
       CALCULATE VOLUME AT NATURAL LENGTH EXCLUDING THE CORE, ft^3
       VN = (PI*(DC**2)*(H1-S))/4.0
       VN = VN+(PI*(DC**2)*(L+S-H1)*(1.0+(D/DC)+(D**2/DC**2)))/12.0
VN = VN-(PI*(DE**2)*L)/4.0
С
     CALCULATE THE VOLUME BELOW EXIT DUCT EXCLUDING THE CORE, ft^3
       VH = (PI*(DC**2)*(H1-S))/4.0
       VH = VH+(PI*(DC**2)*(H-H1)*(1.0+(B/DC)+(B**2/DC**2)))/12.0
       VH = VH - (PI*(DE**2)*(H-S))/4.0
```

C CALCULATE VORTEX EXPONENT

VE = 1.0 - (1.0 - (((12.0 * DC) * * 0.14)/2.5)) * (((T+460.0)/530.0) * * 0.3)

```
C CALCULATE RELAXATION TIME, RT in sec.
```

RT = (DENP*(DPI)**2)/(18.0*VIS)

VAL = S+L

```
IF (VAL .GT. H) THEN VS = VH ELSEIF (VAL .LE. H) THEN VS = VN ENDIF
```

C CALCULATE THE NUMBER OF INLET VELOCITY HEADS, NH

NH = (16.0*A1*B1)/DE**2

C CALCULATE THE INLET VELOCITY, VI:ft/s.

VI = Q/(A1 * B1)

C CALCULATE THE CYCLONE PRESSURE DROP, in H20

DELP = (0.003*DENF*(VI**2)*NH)

C CALCULATE SALTATION VELOCITY, VSA: ft/s.

W = ((4.0*GC*VIS*(DENP-DENF))/(3.0*DENF**2))**(1.0/3.0) VAL1 = B1/DC VAL2 = (VAL1/(1.0-VAL1)**(1.0/3.0)) VSA = 2.055*W*VAL2*(DC**0.067)*(VI**0.667)

C CALCULATE THE OPTIMAL RATIO, VI/VSA

```
RATIO = VI/VSA
```

C CALCULATE CYCLONE CONFIGURATION FACTOR SPECIFIED BY THE C GEOMETRIC RATIOS THAT DESCRIBE THE CYCLONE'S SHAPE

G = (2.0*(PI*(S-(A1/2.0))*((DC**2)-(DE**2)))+4.0*VS)* * (DC/((A1**2)*(B1**2)))

C CALCULATE CYCLONE EFFICIENCY

EFF = 1.0-EXP(-2.0*((G*RT*Q*(VE+1.0))/(DC**3))**(0.5/(VE+1.0))) EFF = 100.0*EFF

C CALCULATE THE EFFECTIVE NUMBER OF TURNS

NT = (0.1079-(0.00077*VI)+(1.924*(10**(-6)))*(VI**2)) NT = NT*VI

с	CALCULATE THE CRITICAL PARTICLE SIZE, ft.
	DPCRIT = (9.0*VIS*DC)/(4.0*PI*NT*VI*(DENP - DENF)) DPCRIT = SQRT (DPCRIT)
с	CALCULATE THE CRITICAL PARTICLE SIZE, microns
	DPMIC = DPCRIT*0.3048*(10**6)
c c	CALCULATE THE RATIO OF THE PARTICLE SIZE TO THE CRITICAL PARTICLE SIZE, RATIO1
	RATIO1 = DPI/DPCRIT
c c	USE THE RESULT TO DETERMINE THE PERCENTAGE REMOVAL OF THE PARTICLES.
	RETURN END
с с с	**************************************
	SUBROUTINE OUTPUT(H,H1,DC,B,A1,B1,S,DE,Q,DENP,DENF,VIS,T,DPI)
	REAL L, NH, NT
	COMMON/RES1/L,D,NH,VI,VSA,RATIO,G,VE,RT,EFF,DELP COMMON/RES2/NT, DPCRIT, DPMIC, RATIO1
100	WRITE (1, 100) FORMAT(///,25x,'CYCLONE DESIGN',/78(1H*))
110 1 2 3	<pre>wRITE (1, 110) H, H1, DC, B FORMAT (5x,'CYCLONE OVERALL HEIGHT, ft:', T60, F9.3,/, 5X,'CYCLONE CYLINDRICAL HEIGHT, ft.:', T60, F9.3,/, 5X,'CYCLONE DIAMETER, ft.:', T60, F9.3,/, 5X,'CYCLONE DUST-OUTLET DIAMETER, ft:', T60, F9.3)</pre>
120 1 2 3	<pre>WRITE (1, 120) A1, B1, S, DE FORMAT (5x,'CYCLONE GAS INLET LENGTH, ft:', T60, F9.3,/, 5x,'CYCLONE GAS INLET WIDTH, ft:', T60, F9.3,/, 5x,'CYCLONE GAS OUTLET LENGTH, ft:', T60, F9.3,/, 5x,'CYCLONE GAS OUTLET DIAMETR, ft:', T60, F9.3)</pre>
130 1 2 3	<pre>WRITE (1, 130) Q, DENP, DENF, VIS FORMAT (5x,'GAS RATE, ft^3/s.:', T60, F10.2,/, 5x,'PARTICLE DENSITY, lb/ft^3:', T60, F9.3,/, 5x,'GAS DENSITY, lb/ft^3:', T60, F9.3,/, 5x,'GAS VISCOSITY, lb/ft.s.:', T60, E12.6)</pre>
	WRITE (1, 140) T, DPI, VE, RT

140 1 2 3	FORMAT (5X, 'GAS TEMPERATURE, oF:', T60, F9.3,/, 5X, 'PARTICLE SIZE (EQUIVALENT DIA.),ft.:',T60,E12.6,/, 5X, 'VORTEX EXPONENT:', T60, F9.4,/, 5X, 'RELAXATION TIME, s:', T60, E12.6)
150 1 2 3	<pre>WRITE (1, 150) VI, VSA, G, DELP FORMAT (5x, 'INLET FLUID VELOCITY, ft/s.:', T60, F9.3,/, 5x,'SALTATION VELOCITY, ft/s.:', T60, F9.3,/, 5x,'CYCLONE CONFIGURATION FACTOR:', T60, F9.3,/, 5x,'CYCLONE PRESSURE DROP, in H20:', T60, F9.3)</pre>
160 1	WRITE (1, 160) RATIO, EFF FORMAT (5X, 'RATIO OF THE INLET TO THE SALTATION VELOCITIES:', T60, F9.3,/,5X,'CYCLONE EFFICIENCY, %:', T60, F9.3)
170 1 2 3	<pre>WRITE (1, 170) NT, DPCRIT, DPMIC, RATIO1 FORMAT (5X,'EFFECTIVE NUMBER OF TURNS:', T60, F8.2,/, 5X,'CRITICAL PARTICLE DIAMETER, ft.:', T60, E14.6,/, 5X,'CRITICAL PARTICLE DIAMETER, microns:',T60,E14.6,/, 5X,'RATIO OF PARTICLE DIA. TO CRITICAL DIA.:',T60,F8.2)</pre>
180	WRITE (1, 180) Format (78(1H-))
с	FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR(12)
	RETURN END

PROGRAM PROG45

~			*********		
с					
с			GAS DRYER.		
С			LL PROVIDE A PRELIMINARY DESIGN FOR A SOLID-		
С			USED FOR THE REMOVAL OF MOISTURE FROM GAS STREAMS.		
c			L CALCULATE WATER PICKUP, DESICCANT VOLUME, DESICCANT		
С	BED (TOWER) DIAMETER, GAS VELOCITY THROUGH THE BED, VESSEL WEIGHT,				
С			ION REQUIREMENTS AND MOMENTUM FOR MAXIMUM DOWNFLOW		
с	VELOCI				
с			***************		
С	H2OL	=	TOTAL WATER ADSORBED, 1b/hr.		
с	FLOW		GAS FLOW RATE, 10^6 std ft^3/day.		
С	CYCLE		DRYING CYCLE, hr.		
с	H2OC		WATER CONTENT OF GAS, 1b/10^6 std ft^3.		
С	DESVOL	=	DESICCANT VOLUME, ft^3.		
с		=	DESICCANT DENSITY, 1b/ft^3.		
С	AREA1	=	CALCULATED AREA BASED ON DESICCANT VOLUME, ft^3.		
С	BL	-	DESICCANT BED LENGTH, ft.		
С		=	CALCULATED BED DIAMETER USING AREA1, ft.		
С	AREA	-	ACTUAL DESSIGANT BED AREA USING BD CALCULATED OR		
с			MODIFIED BY THE USER, ft^2.		
С	Q	=	ACTUAL GAS FLOW RATE AT FLOWING CONDITIONS, ft^3/min.		
С	т	=	GAS FLOWING TEMPERATURE OF.		
С	Р	=	DRYER OPERATING PRESSURE, psig.		
с	Z	-	COMPRESSIBILITY FACTOR		
с	VEL	2	GAS VELOCITY, ft/min.		
с	TC	=	DRYER SHELL THICKNESS, in.		
С	н	=	ADDITIONAL SHELL HEIGHT FOR DESICCANT SUPPORTS		
с			AND DISTRIBUTOR, ft.		
С	R	=	RADIUS OF DESICCANT BED, ft.		
с	S	-	ALLOWABLE STRESS, 1b/in^2.		
с	Е	=	JOINT EFFICIENCY.		
с	THEAT	=	TOTAL HEAT REQUIRED TO REGENERATE DRYER (IDEAL), Btu.		
с		=	STEEL WEIGHT OF SHELL AND HEADS, 1b.		
с	WD	=	DESICCANT WEIGHT, 1b.		
с	тв	-	TEMPERATURE OF DESICCANT BED, oF.		
c	нт	=	HEATER TEMPERATURE, OF.		
c	F	=	LOSS FACTOR FOR NON-STEADY STATE HEATING.		
c	REGENH	-	HEAT ADDED TO REGENERATION GAS TO REGENERATE		
c			DESICCANT BED.		
c	H1,H2	=	ENTHALPY OF REGENERATION GAS BEFORE		
с			AND AFTER REGENERATION, Btu/lb		
c	REGENG	=	TOTAL REGENERATION GAS REQUIREMENT, 1b		
c	MW	=	MOLECULAR WEIGHT OF REGENERATION GAS		
č	SCF	_	STANDARD CUBIC FEET OF REGENERATION GAS		
č	PICKUP		DESICCANT WATER CAPACITY, %		
č			***************************************		

REAL MW, MOMEN

COMMON/DRY1/ FLOW, P, T COMMON/DRY2/ Z, CYCLE, H2OC COMMON/DRY3/ S, E, PICKUP COMMON/DRY3/ DDEN, BL, HT COMMON/DRY5/ MW, H1, H2

COMMON/RES1/ H2OL, DESVOL, AREA1 COMMON/RES2/ BD, R, Q, VEL

COMMON/RES3/ TC, VESWT, WD COMMON/RES4/ TB, THEAT, F COMMON/RES5/ REH, REG, SCF, MOMEN OPEN (UNIT=3, FILE='DATA45.DAT', STATUS='OLD', ERR=18) OPEN (UNIT=1, FILE='PRN') C C C READ GAS FLOW RATE, FLOW: 10^6 std ft^3/day. READ DRYER OPERATING PRESSURE, P: psig. READ GAS FLOWING TEMPERATURE, T: oF. READ GAS COMPRESSIBILITY FACTOR, Z. READ DRYING CYCLE, Nr. READ DRYING CYCLE, CYCLE: hr. READ WATER COMTENT OF GAS, H2OC: 1b/MM std ft^3. READ ALLOWABLE STRESS, S: 1b/in^2. READ JOINT EFFICIENCY, E: 1.0 READ DESICCANT WATER CAPACITY, PICKUP: \$ READ DESCICCANT DENSITY, DDEN: 1b/ft^3. READ DESICCANT BED LENGTH, BL: ft. READ HEATER TEMPERATURE, HT: OF. READ MOLECULAR WEIGHT OF REGENERATING GAS, MW: READ ENTHALPY OF REGENERATION GAS BEFORE REGENERATION HEATER, H1: Btu/lb. READ ENTHAPLY OF REGENERATION GAS AFTER REGENERATAION HEATER, H2: Btu/lb. READ (3, *, ERR=19) FLOW, P, T READ (3, *, ERR=19) Z, CYCLE, H2OC READ (3, *, ERR=19) S, E, PICKUP READ (3, *, ERR=19) DDEN, BL, HT READ (3, *, ERR=19) MW, H1, H2 GO TO 10 WRITE (*, 11) 18 FORMAT (6X, 'ERROR MESSAGE IN THE DATA VALUE') 11 GO TO 999 WRITE (*, 12) FORMAT (6X, 'DATA FILE DOES NOT EXIST') 19 12 GO TO 999 с CALCULATE THE REQUIREMENTS FOR A GAS DRYER CALL DRYER 10 с PRINT THE RESULTS. CALL OUTPUT CLOSE (UNIT=3, STATUS='KEEP') CLOSE (1) 999 STOP END с C C C THIS PROGRAM SIZES THE GAS DRYER WITH THE GIVEN DATA VALUES

SUBROUTINE DRYER PARAMETER (PI=3.1415927) REAL MW, MOMEN COMMON/DRY1/ FLOW, P, T COMMON/DRY2/ Z, CYCLE, H2OC COMMON/DRY3/ S, E, PICKUP COMMON/DRY3/ DEEN, BL, HT COMMON/DRY5/ MW, H1, H2 COMMON/RES1/ H2OL, DESVOL, AREA1 COMMON/RES2/ BD, R, Q, VEL COMMON/RES3/ TC, VESWT, WD COMMON/RES4/ TB, THEAT, F COMMON/RES5/ REH, REG, SCF, MOMEN с CALCULATE TOTAL WATER ADSORBED, H2OL. H2OL = (FLOW*CYCLE*H2OC)/24.0с CALCULATE DESICCANT VOLUME, DESVOL DESVOL = (H2OL*100.0)/(PICKUP*DDEN) с CALCULATE DESICCANT CROSS-SECTIONAL AREA, AREA1. AREA1 = DESVOL/BL с CALCULATE BED DIAMETER, BD. BD = SQRT((4.0 * AREA1)/PI)с CALCULATE THE BED AREA, AREA. AREA = (PI*BD**2)/4. T1 = T+460.R = (12.*BD)/2.P1 = P+14.7 CALCULATE THE ACTUAL GAS FLOW RATE AT FLOWING CONDITIONS, ft^3/min. С Q = (FLOW * T1 * Z * 19.6314)/P1С CALCULATE THE SUPERFICIAL GAS VELOCITY, ft./min. VEL = Q/AREA CALCULATE THE DRYER SHELL THICKNESS, in. С TC = (P*R)/(S*E-0.6*P)

C ASSUME H=2.0 ft. ADDITIONAL SHELL HEIGHT FOR DESICCANT SUPPORT AND

```
C DISTRIBUTOR.
```

H = 2.0

R1 = BD/2.0

C CALCULATE THE VESSEL WEIGHT., 1b.

```
VESWT = (480.*PI*BD*(TC/12.0)*(BL+H+R1)*1.1)
TB = 0.75*HT+0.25*(HT-50.)
```

WD = DDEN*DESVOL

THEAT = 1.05*(VESWT*0.12*(TB-T)+WD*0.25*(TB-T)+H2OL*1400.) F = ALOG((HT-T)/50.)

```
C CALCULATE TOTAL REGENERATION GAS REQUIREMENT AND STANDARD CUBIC
C FEET OF REGENERATION GAS
```

REH = THEAT*F REG = REH/(H1-H2) SCF = (REG*380.0)/MW

```
C CALCULATE ALCOA'S MOMENTUM FOR MAXIMUM DOWNFLOW
C VELOCITIES i.e. DESICCANT ATTRITION SHOULD NOT BE A PROBLEM
C IF THE MOMENTUM IS EQUAL TO OR LESS THAN 30,000.
C MOMENTUM = (V)(MOL.WT) (P. Atms)
```

MOMEN = VEL*MW*(P1/14.7)

RETURN END

SUBROUTINE OUTPUT

REAL MOMEN, MW

COMMON/DRY1/ FLOW, P, T COMMON/DRY2/ Z, CYCLE, H2OC COMMON/DRY3/ S, E, PICKUP COMMON/DRY3/ DDEN, BL, HT COMMON/DRY5/ MW, H1, H2

COMMON/RES1/ H2OL, DESVOL, AREA1 COMMON/RES2/ BD, R, Q, VEL COMMON/RES3/ TC, VESWT, WD COMMON/RES4/ TB, THEAT, F COMMON/RES5/ REH, REG, SCF, MOMEN

с

```
WRITE (1, 100)
100 FORMAT (///,35X, 'GAS DRYER DESIGN',/1H ,78(1H*))
```

110	*	<pre>wRITE (1, 110) FLOW, P, T FORMAT (5X, 'GAS FLOW RATE, 10^6 std. ft^3/day:', T60, F12.3,/, 5X,'DRYER OPERATING PRESSURE, psig.:', T60, F12.3,/, 5X,'GAS FLOWING TEMPERATURE, oF:', T60, F12.1)</pre>
120	*	<pre>WRITE (1, 120) Z, CYCLE, H2OC FORMAT (5X,'GAS COMPRESSIBILITY FACTOR:', T60, F12.4,/, 5X,'DRYING CYCLE, hr.:', T60, F12.1,/, 5X,'WATER CONTENT OF GAS, lb/10^6 std.ft^3:',T60,F12.0)</pre>
130	*	<pre>WRITE (1, 130) S, E, PICKUP FORMAT (5X,'ALLOWABLE STRESS, psi:', T60, F12.1,/, 5X,'JOINT EFFICIENCY:', T60, F12.1,/, 5X,'USEFUL DESIGN CAPACITY OF DESICCANT, (%):', T60, F12.1)</pre>
140	*	WRITE (1, 140) DDEN, HT FORMAT (5X, 'DESICCANT DENSITY, 1b/ft^3:', T60, F12.1,/, 5X, 'HEATER OUTLET TEMPERATURE, oF:', T60, F12.1)
150	*5 *F	WRITE (1, 150) MW, H1, H2 FORMAT (5X, 'MOLECULAR WEIGHT OF REGENERATION GAS:',T60,F12.1,/, X,'ENTHALPY OF GAS BEFORE REGENERATION HEATER, Btu/lb:',T60, 12.1,/,5X,'ENTHALPY OF GAS AFTER REGENERATION HEATER, Btu/lb:', 60, F12.1)
160	*	WRITE (1, 160) H2OL, DESVOL, AREA1 FORMAT (5X,'TOTAL WATER ADSORBED, 1b/h.:', T60, F12.3,/, 5X,'DESICCANT VOLUME, ft^3.:', T60, F12.4,/, 5X,'CALCULATED CROSS-SECTIONAL AREA, ft^2.:', T60, F12.4)
170	* * *	<pre>WRITE (1, 170) BD, R, BL, Q, VEL FORMAT (5X,'BED DIAMETER, ft.:', T60, F8.4,/, 5X,'BED RADIUS, ft.:', T60, F8.4,/, 5X,'BED LENGTH, ft.:', T60, F8.2,/, 5X,'ACTUAL GAS FLOW RATE, ft-3/min.:', T60, F12.3,/, 5X,'SUPERFICIAL GAS VELOCITY, ft./min.:', T60, F12.3)</pre>
180	*	<pre>WRITE (1, 180) TC, VESWT, WD FORMAT (5X,'DRYER SHELL THICKNESS, in.:', T60, F12.4,/, 5X,'VESSEL WEIGHT, lb.:', T60, F12.0,/, 5X,'DESICCANT WEIGHT, lb.:', T60, F12.0)</pre>
190	* *	<pre>WRITE (1, 190) TB, THEAT, F FORMAT (5X,'TEMPERATURE OF DESICCANT BED, oF:', T60, F12.3,/, 5X,'TOTAL HEAT REQUIRED TO REGENERATE DRYER, Btu:', T60, E14.6,/, 5X, 'LOSS FACTOR:', T60, F12.4)</pre>
200	* * *	<pre>wRITE (1, 200) REH, REG, SCF, MOMEN FORMAT (5X,'HEAT ADDED TO REGENERATION GAS:', T60, E14.6,/, 5X,'TOTAL REGENERATION GAS REQUIREMENT, lb:',T60, E14.6, /,5X,'STANDARD CUBIC FEET OF REGENERATION GAS, std.ft^3:', T60, E14.6,/,5X,'MOMENTUM:', T60, E14.6)</pre>
210		WRITE (1, 210) FORMAT (76(1H-))
с		FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.
		WRITE (1, *) CHAR(12)
		RETURN END

CHAPTER 5

Instrument Sizing

INTRODUCTION

Instruments are used to monitor the pertinent process variables during plant operation. They may be part of automatic control loops or used for the manual monitoring of the process operation. Instruments that monitor critical process variables are fitted with automatic alarms to alert the operators to critical and hazardous situations. Miller [1] and Perry [2] provide details of process instruments and control equipment.

This chapter considers orifice sizing for liquid and gas flows, control valve design for liquid, vapor, steam services, and two-phase flow. The chapter further reviews relief valve sizing for vapor, steam, liquid, air, and fire conditions. The effect of the pressure drop (ΔP) on pipework design, noise suppression, discharge reactive forces, and relief valve location is considered. In addition, the chapter discusses two-phase flow relief sizing involving runaway reactions. For all the sizing procedures, the chapter develops the engineering equations for ease in preparing computer programs.

Orifice Sizing For Liquid and Gas Flows

In process design, an orifice or a nozzle is usually sized for known flow rate and pressure drop (ΔP). Nozzles and orifices are used to restrict the flow of fluids. Thus, a reduction in cross-sectional area can cause significant changes in the pressure and temperature of a compressible fluid as it flows through the constriction. The magnitude of the change depends on the linear velocity of the fluid or the Mach number at the constriction. An orifice is usually designed to develop a full-scale pressure differential at maximum flow. Alternatively, it can be designed to a square-edged or sharp-edged cut. A square-edged or sharp-edged orifice is a clean cut square-edge hole with straight walls perpendicular to the flat upstream face of a thin plate placed crosswise of the channel. The stream issuing from such an orifice attains its minimum cross section (vena-contracta) at a distance downstream of the orifice. This varies with the ratio, β , of the orifice to the pipe diameter. All measurements of the distance from the orifice are made from the upstream face of the plate for the corner, radius, pipe, and vena contracta taps.

Corner Taps. Static holes are drilled, one in the upstream and the other in the downstream flange. The opening is as close as possible to the orifice plate.

Radius Taps. Static holes are located one pipe diameter upstream and one-half diameter downstream from the plate.

Pipe Taps. Static holes are located $2\frac{1}{2}$ pipe diameters upstream and 8 pipe diameters downstream from the plate.

Vena-Contracta Taps. The upstream static hole is $\frac{1}{2}$ to 2 pipe diameters from the plate; the downstream tap is located at the position of minimum pressure.

Flange Taps. Static holes are located 1 inch upstream and 1 inch downstream of the plate.

Figure 5-1 shows the locations and names of the orifice taps. The true mass flow rate for both liquids and gases is obtained by multiplying the theoretical mass flow equation by the discharge coefficient, the gas expansion factor, and the thermal-expansion factor. This is expressed as:

$$W_{true} = 0.52502 \left(\frac{C_{\rm D} Y d^2 F_{\rm a}}{\sqrt{1 - \beta^4}} \right) (\rho \Delta P)^{0.5}$$
(5-1)

where the coefficient of discharge, C_D , is

$$C_{\rm D} = \frac{\text{true flow rate}}{\text{theoretical flow rate}}$$
(5-2)

 C_D for a given orifice type is a function of the Reynolds number (N_{Re}) and the diameter ratio β . C_D is constant at Reynolds number greater than 30,000. For square-edged or sharp-edged concentric circular

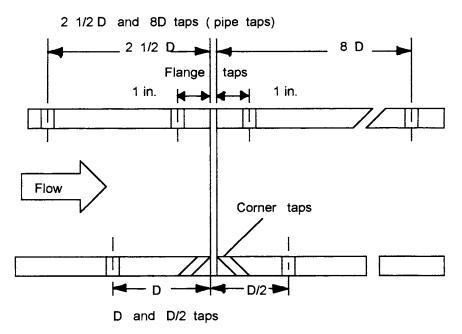


Figure 5-1. Orifice pressure tap locations. 2 1/2D and 8D pipe taps are not recommended in ISO 5167 or ASME fluid meters. D and D/2 taps are now used in place of vena-contracta taps.

orifices, C_D falls between 0.595 and 0.62. The discharge coefficient corrects the theoretical equation for the influence of velocity profile (N_{Re}), assuming no energy loss between the taps and pressure-tap location.

The thermal expansion factor F_a : The material of primary element and pipe expands or contracts with temperature. The pipe and bore diameters are measured at room temperature, but will be larger or smaller when used at other temperatures. F_a is introduced to correct for these differences.

The Expansion Factor Y. This corrects for density changes between taps. Y depends on the adiabatic exponent ($k = C_p/C_v$), absolute pressure ratio ($\Delta P/P_1$), and the ratio, β , between restriction diameter and the inside pipe diameter. For liquid flow, Y = 1.0. For gas flow, Buckingham derived the equation [3]:

$$Y = 1 - (0.41 + 0.35\beta^4) \frac{\Delta P}{Pk}$$
(5-3)

Orifice Sizing For Liquid Flows

For liquid sizing, Y=1.0, and the following assumptions are $F_a = 1.0$, $C_D = 0.62$ [4]

Equation 5-1 is re-arranged in the form

$$\frac{\sqrt{1-\beta^4}}{d^2} = \frac{0.52502(C_D)}{W_{true}} (\rho \Delta P)^{0.5}$$
(5-4)

where

$$Z = \frac{0.52502(C_{\rm D})}{W_{\rm true}} (\rho \Delta P)^{0.5}$$
(5-5)

and

$$\beta = \frac{d}{D}$$
(5-6)

Equation 5-4 is expressed by

$$\frac{1}{d^2} \left[1 - \left(\frac{d}{D}\right)^4 \right]^{0.5} = Z$$
(5-7)

squaring both sides of Equation 5-7

$$\frac{1}{d^4} \left[1 - \left(\frac{d}{D}\right)^4 \right] = Z^2$$
(5-8)

$$\frac{1}{d^4} - \frac{1}{D^4} = Z^2 \tag{5-9}$$

$$\frac{1}{d^4} = Z^2 + \frac{1}{D^4}$$
(5-10)

The orifice size, d, is expressed by

$$d = \left(\frac{D^4}{1 + D^4 Z^2}\right)^{0.25}$$
(5-11)

Orifice Sizing For Gas Flows

Using Equations 5-1 and 5-3 That is,

$$W_{true} = 0.52502 \left(\frac{C_{\rm D} Y d^2 F_{\rm a}}{\sqrt{1 - \beta^4}} \right) (\rho \Delta P)^{0.5}$$
(5-1)

and

$$Y = 1 - (0.41 + 0.35\beta^4) \frac{\Delta P}{Pk}$$
(5-3)

The following substitutions are made:

$$y = \beta^4 \tag{5-12}$$

$$1 = \frac{W_{true}}{0.52502(D^2)(\rho\Delta P)^{0.5}}$$
(5-13)

$$n = 1 - 0.41 \frac{\Delta P}{P.k}$$
(5-14)

$$t = 0.35 \frac{\Delta P}{P.k}$$
(5-15)

Equation 5-1 and Equation 5-3 are modified and combined to give

$$l = \frac{(ny^{1/2} - ty^{3/2})}{(1 - y)^{1/2}}$$
(5-16)

Squaring both sides and dividing by t², Equation 5-16 is expressed in cubic form as:

$$y^{3} - \frac{2n \cdot y^{2}}{t} + \frac{n^{2} + l^{2}}{t^{2}}y - \frac{l^{2}}{t^{2}} = 0$$
(5-17)

Equation 5-17 is solved for y by finding roots of the cubic equation using the following substitutions.

Let

$$p = \frac{2n}{t} \tag{5-18}$$

$$q = \left(\frac{n^2 + l^2}{t^2}\right) \tag{5-19}$$

$$r = -\frac{l^2}{t^2}$$
(5-20)

$$y^3 - 2py^2 + qy - r = 0 (5-21)$$

and

$$a = \frac{3q - p^{2}}{3}$$
$$= \frac{3l^{2} - n^{2}}{3t^{2}}$$
(5-22)

$$b = \frac{(2p^{3} - 9pq + 27r)}{27}$$
$$= -\frac{(2n^{3} + 18nl^{2} + 27tl^{2})}{27t^{3}}$$
(5-23)

The solution to Equation 5-17 yielding a real number is:

$$y = \left\{ -\frac{b}{2} + \left(\frac{b^2}{4} + \frac{a^3}{27}\right)^{1/2} \right\}^{1/3} + \left\{ -\frac{b}{2} - \left(\frac{b^2}{4} + \frac{a^3}{27}\right)^{1/2} \right\}^{1/3} - \frac{2n}{3t}$$
(5-24)

the orifice diameter, d, is expressed by

$$d = Dy^{1/4}$$
 (5-25)

Conversions:

 $\Delta P = 0.03613(h_w)$ (5-26)

$$W_{true} = \left(\frac{Q.\rho}{60.0}\right)$$
(5-27)

The orifice sizing applies to corner, flange, vena-contracta and 1/2diameter taps.

Nomenclature

 $C_{\rm D}$ = coefficient of discharge

- d = orifice diameter for full scale ΔP , inch
- D = internal pipe diameter, inch
- F_a = area thermal-expansion factor. Takes into account the expansion of the orifice with temperature changes
- $h_w =$ full-scale orifice pressure differential, in. H₂O at 68°F
- k = specific heat ratio (constant pressure to constant volume C_p/C_v)

P = upstream pressure, psia

 ΔP = full-scale orifice pressure-differential, psi

Q = full-scale flow rate at upstream conditions, ft³/min

 W_{true} = true full-scale flow rate, lb/s

Y = expansion factor

 β = d/D, ratio of orifice diameter to pipe I.D.

 ρ = upstream fluid density, lb/ft³

CONTROL VALVE SIZING

Introduction

Valves and piping represent about 25 percent of the total capital expenditure for materials and equipment in the chemical process industries (CPI). These industries employ valves in a wide range of flowing media from granula solids to industrial wastes. The most important characteristics in the selection of valves are viscosity, corrosiveness, and abrasiveness. In addition, a designer must consider the process parameters. These may be predictable short-term upset conditions, handling more than one fluid with the same valve, or potentially high pressures resulting from a fluid being entrapped in a valve body that then vaporizes due to heat gain. Selecting the right control valve for an application involves four basic styles of throttling: cage-style globe valves, ball valves, eccentric-disk valves, and butterfly valves. Globe valves are the standards of the control valves. These may be hand operated or power actuated types for automatic control. The common feature of these valves is their internal construction, which consists of a disc or plug reciprocating within the valve body. Figure 5-2A shows a typical globe valve, and Figure 5-2B shows a globe valve with cage-style trim.

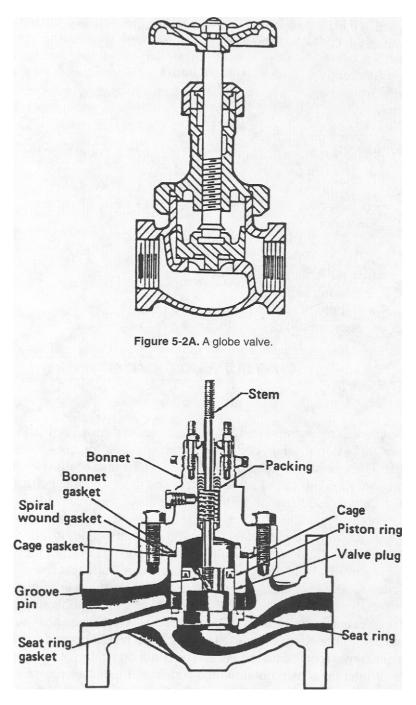


Figure 5-2B. A globe valve with cage-style trim.

Langford [5] has reviewed the installation, maintenance and troubleshooting of control valves, while Driskell [6] studied the various aspects of two-phase flow, non-turbulent, non-Newtonian and flashing liquids. Recently, Collins [7] listed the main basic conditions for selecting control valves. These include (1) characteristics of the controlled fluid, (2) system pressures, (3) maximum differential pressure, (4) fluid operating temperature and range, (5) line size, (6) valve size, (7) valve type, (8) valve C_v , (9) allowable leakage, (10) end connections, (11) valve environment (outside temperature and range), (12) seat material, body material, (13) actuation medium, (14) flow characteristics, and (15) rangeability.

Correctly sized control valves achieve a high quality of control. A valve that is too small will not allow the required flow, and a valve that is too large will cost more than a correctly sized smaller valve and the flow will be uncontrollable because its full control range will not be used. The procedure for sizing a control valve is [8]:

- 1. Calculate the required valve sizing coefficient, C_v, based on process data and manufacturers' valve data.
- 2. Consult valve manufacturers' table of C_v against size. Then select the smallest valve with a C_v rating greater than or equal to the required C_v .
- 3. Check that the reducers required installing the valve will not change the valve selection. Choose a new valve with a greater C_v if necessary.
- 4. Check that the flow through the valve is not choked.

When the liquid flow is turbulent, and the effects of choked flow and reducers are ignored, the expression for C_v reduces to

$$C_{v} = \frac{q}{K_{D}} \left(\frac{SpGr_{f}}{\Delta P}\right)^{0.5}$$
(5-28)

Pressure Drop For Sizing

The pressure drop (ΔP_{sizing}) used for sizing the control valve is the actual pressure drop across the valve at the sizing flow rate. This is obtained by starting with the system pressure drop (ΔP_{system}) . The pressure drop at the sizing flow rate of all pipe, fittings, and equipment in the flow system is subtracted from ΔP_{system} . The remainder is the actual pressure drop across the control valve at the sizing flow rate.

Choked Flow

As the flow of liquid passes the point of greatest restriction inside the control valve, its velocity reaches a maximum and its pressure falls to a minimum. If the pressure falls below the vapor pressure of the liquid, vapor bubbles form within the valve. Increasing ΔP_{sizing} across the valve beyond the point where vapor bubbles form would not alter the flow. This suggests choked flow. The ΔP at which choked flow begins is called the terminal, ΔP_{T} . ΔP_{T} is used instead for sizing the control valve when ΔP_{sizing} across the valve is greater than ΔP_{T} .

Flashing and Cavitation

Choked flow produces either flashing or cavitation. Flashing occurs when the pressure downstream of the valve is below the vapor pressure of the liquid, ΔP_v . Pipe and internal valve erosion often occur because the velocity of flashing the vapor-liquid stream is higher than the inlet liquid velocity. A cavitation will occur if the downstream pressure is above the liquid's vapor pressure. This results in the collapse of the vapor bubbles as they leave the point of greatest restriction in the valve. The shock waves and noise caused by the collapsing bubbles cause rapid and severe damage to the valve or piping.

CONTROL VALVE SIZING FOR LIQUID, GAS, STEAM, AND TWO-PHASE FLOW

The valve flow coefficients, C_v for a given valve depend on the internal dimensions of the valve and the smoothness of the valve's internal surface. Control valve manufacturers establish the C_v for their valves through a series of tests using water or air at a given pressure differential across the valve. Manufacturers define C_v as the capacity index. This indicates the flow of water (gpm at 60°F) that will pass through a completely open valve under a pressure differential between the inlet and outlet flanges. Control valve coefficients for single and double seated valves are given in Table 5-1 [9].

Liquid Sizing

The equation for sizing valve for liquid service is given by

$$C_v = Q_1 \left(\frac{SpGr_1}{P_1 - P_2}\right)^{0.5}$$
 (5-29)

Size, in.	Flow Coefficient Single-Seat	C _v Double-Seat
3/4		8
1	9	12
1 1/4	14	18
1 1/2	21	28
2	36	48
2 1/2	54	72
3	75	110
4	124	195
6	270	450
8	480	750
10	750	1160
12	1080	1620
14	1470	2000
16	1920	2560

Table 5-1Flow Coefficients for Control Values

Alternatively, for a given C_v , the liquid flow rate can be defined by

$$Q_{1} = C_{v} \left(\frac{P_{1} - P_{2}}{SpGr_{1}}\right)^{0.5}$$
(5-30)

Equation 5-29 is valid for liquid flowing below its saturation temperature in the turbulent zone with a viscosity value that is close to that of water and size of pipe. Also, the control valve must be the same. Equation 5-29 can also be applied, if the vapor pressure of the liquid at the flowing temperature is equal to or less than one-half the upstream pressure. For this case, the vapor pressure of the liquid is substituted for downstream pressure, $P_{2,}$ and the valve coefficient is calculated. The calculated C_v must be corrected by a critical flow factor, C_f , where C_v corr is defined by

$$C_v \text{corr} = \frac{C_v \text{calc}}{C_f}$$
(5-31)

The value of C_f is between 0.85 and 0.98. Flow capacity of a control valve placed between pipe reducers is slightly decreased. In subcritical

flow, this is accounted for by a correction factor, R. In critical flow, the correction factor is C_{fr} . This replaces C_f in the sizing calculations. In addition, R, C_f , and C_{fr} depend on the ratio between pipe size and valve size. Table 5-2 lists values of R, C_f , and C_{fr} for single-seated and double-seated control valves [9].

Gas Sizing

The following equations are used for sizing control valves for gas and steam conditions. These equations apply to both subcritical and critical flow conditions [10].

For subcritical condition, C_v , is defined by

$$C_{v} = \frac{Q_{g} \{ (SpGr_{g})(T) \}^{0.5}}{1360 \{ (P_{1} - P_{2})(P_{2}) \}^{0.5}}$$
(5-32)

		Single	Seat	Double	Seat
Condition	Factor	Equal- Percentage	V-Port	Equal- Percentage	V-Port
Critical flow line size control valve	C _f	0.98 or .85	0.98	0.98	0.98
Critical flow (control valve between pipe reducers)	C _{fr}	0.86	0.94	0.86	0.94
Subcritical flow $D/d = 1.5$	R	0.96	0.96	0.96	0.94
Subcritical flow D/d = 2 (Control valve between pipe reducers)	R	0.94	0.94	0.94	0.94

Table 5-2 Correction Factors for Control Valve Flow Coefficient

The gas flow rate is computed for a given C_v , and is defined by

$$Q_{g} = \frac{C_{v}(1360) \{ (P_{1} - P_{2})(P_{2}) \}^{0.5}}{\{ (SpGr_{g})(T) \}^{0.5}}$$
(5-33)

Critical Condition

For critical flow, the downstream pressure of the control valve is less than one-half the upstream pressure. Therefore, the term $\{(P_1 - P_2)(P_2)\}^{0.5}$ reduces to $0.5P_1$ in Equations 5-32 and 5-33. C_v is then defined by

$$C_{v} = \frac{Q_{g} \{ (SpGr_{g})(T) \}^{0.5}}{1360(0.5P_{1})}$$
(5-34)

The gas flow rate is:

$$Q_{g} = \frac{C_{v}(1360)(0.5P_{1})}{\left\{(SpGr_{g})(T)\right\}^{0.5}}$$
(5-35)

Steam Sizing

The following equations are used to size control valves for saturated and superheated steam flow. For saturated steam flow, a value of zero is used for °F superheat.

Subcritical Condition. The control valve coefficient, C_v , is defined by

$$C_{v} = \frac{(W)(K)}{3\{(P_{1} - P_{2})(P_{2})\}^{0.5}}$$
(5-36)

The steam flow rate, W, is defined by

$$W = \frac{(3)(C_v) \{ (P_1 - P_2)(P_2) \}^{0.5}}{K}$$
(5-37)

Critical Condition. The control valve coefficient and the steam flow rate are expressed as:

$$C_{v} = \frac{(W)(K)}{1.5P_{1}}$$
(5-38)

and

$$W = \frac{1.5(P_{\perp})(C_{\nu})}{K}$$
(5-39)

where

$$K = 1 + (0.0007 \times {}^{\circ}F \text{ superheat})$$
 (5-40)

Two-Phase Flow

Sizing a control valve for liquid-gas mixtures introduces incorrect valve coefficient, C_v , if C_v for each of the two fractions is calculated separately, and then added. This method assumes that the liquid and gas are passing through the valve orifice independently and at greater different velocities.

Driskell [6] has given a plausible assumption that the two fluids travel through the throttling orifice as a homogeneous mixture at approximately the same velocity. He applied the gas expansion factor to the gas fraction only, and not to the total mixture. C_v for liquid-gas mixtures can be expressed as follows:

Divide the specific volume of the inlet gas, v_g , by the square of the expansion factor Y. This gives an effective incompressible specific volume, while the liquid specific volume, v_l , is unaffected.

That is,

$$\frac{v_g}{Y^2} + v_1 \tag{5-41}$$

where $v_g = 1/\rho_g$ and $v_l = 1/\rho_l$

Add the effective volumes of the two fluids in proportion to their mass fraction, f, in the mixture to obtain the effective specific volume of the stream, v_e .

$$\mathbf{v}_{e} = \frac{\mathbf{f}_{g} \cdot \mathbf{v}_{g}}{\mathbf{Y}^{2}} + \mathbf{f}_{1} \cdot \mathbf{v}_{1}$$
 (5-42)

where the expansion factor is:

$$Y = 1 - \frac{x}{(3F_k x_T)}$$
(5-43)

and

 $\begin{aligned} \mathbf{x} &= \Delta P/P_{i} \\ \mathbf{x}_{T} &= \text{pressure drop ratio factor (from manufacturer)} \\ \mathbf{F}_{k} &= \text{ratio of specific heats factor (k/1.40)} \\ \mathbf{k} &= \text{ratio of specific heats (} \mathbf{C}_{p}/\mathbf{C}_{v}) \\ \Delta P &= P_{1} - P_{2}, \text{psi} \\ P_{1} &= \text{inlet pressure, psia} \\ P_{2} &= \text{outlet pressure, psia} \end{aligned}$

The valve coefficient, C_v , is:

$$C_{v} = \frac{W}{63.3 \left(\frac{\Delta P}{v_{e}}\right)^{0.5}}$$
(5-44)

Alternatively, for a given C_v , the flow rate in lb/h is:

$$W = 63.3(C_v) \left(\frac{\Delta P}{v_e}\right)^{0.5}$$
(5-45)

Installation

Control valves form an important equipment item in process plants, and therefore, a lot of money can be saved by ensuring that the right valves are installed. Selecting the right control valve for any application starts with defining the valve's functions. For example, if an on-off interlock valve is to be used, then the primary criteria for selection are reliability and simplicity. If very fast and precise modulation is required, such as for a compressor antisurge vent, a high quality valve is chosen.

Factors that affect the selection of control valves are:

- **Defining the type of fluid.** This includes the physical properties (density and viscosity), corrosive properties, and the nominal pressures, temperatures and flow rate. In the case of fluids, it is necessary to know the vapor pressures to check for flashing and cavitation. Also a high temperature increase can severely damage some types of gaskets and packings.
- *The valve size*. This can be defined using standard calculation methods. The operating conditions and calculations may eliminate cer-

tain valve styles from consideration. Cavitation is more evident with the high-recovery rotary type valves and sometimes can be avoided by selecting a globe valve. In addition, a valve that is too small will not permit the required amount of flow. Alternatively, a valve that is too large will be expensive and may create trim wear and control problems.

- Access for maintenance. This must be provided during the design phase, retained during construction, and considered during the revision stage. Maintenance can be both problematic and expensive if a valve is difficult to reach or to work on. A substantial change in the fluid-momentum (e.g., a flow through an angle valve or a large fluid-velocity change) causes high reaction forces. This can cause serious damage to the valves. The valve should be installed between one and three feet above the grade or platform to control the process.
- *Excessive stress on a valve body.* This can cause leakage and prevent normal operation of the moving parts resulting in the breakage of the flange or valve body. Poor pipe alignment is the most probable cause of excess stress. Also, heavy valves require support in order to reduce the stress and to make removal and reinstallation easier.

Noise

Cavitation refers to the formation and subsequent collapse of vapor bubbles in the flowing liquid stream at a point of higher pressure. This causes erosion of the valve parts. It generates extremely high pressure shock waves that hammer against the valve outlet and piping. Pressures in these collapsing cavities can be as high as 500,000 psi in magnitude, and often result in severe and rapid damage to the valve and piping.

Noise from control valves is often caused by pressures, flows, and temperatures involving compressible flow. Valves control flow by causing turbulence, thereby converting much of the energy to heat and a small fraction to sound. The control valve manufacturers have developed noise reducing valve trims and high-capacity valve configuration. Langford [5] suggested that noise may be reduced by as much as 20 dB by installing acoustic insulation at the valve and downstream, or alternatively by using heavier wall pipe downstream.

Nomenclature

- $C_v = capacity \text{ coefficient for fully open control valve}$
- f_g = mass fraction in the gas phase

 $f_1 = mass$ fraction in the liquid phase F_k = ratio of specific heats factor (k/1.40) k = ratio of specific heats (C_p/C_v) K = superheat correction factor $K_{\rm D}$ = coefficient of discharge P_1 = inlet pressure, psia P_2 = outlet pressure, psia $\Delta P = pressure drop, psi$ q = fluid flow rate Q_1 = liquid flow rate, gpm Q_g = gas flow rate,ft³/h at 14.7 psia and 60°F $SpGr_{f}$ = specific gravity of fluid $SpGr_1 = specific gravity of liquid$ $SpGr_a = specific gravity of gas = M Wt of gas/M Wt of air$ \overline{T} = flowing temperature, °R (°F + 460) V_{e} = effective specific volume of the stream V_{α} = specific volume of the inlet gas V_1 = specific volume of the inlet liquid X_{T} = pressure drop ratio factor (from manufacturer) W =steam flow, lb/h

RELIEF VALVE SIZING

Introduction

Safety relief valves serve a valuable function in protecting against overpressure of pipework and process equipment. This may occur due to operational malfunction, such as equipment failure, fire or human error. Proper sizing of a relief system is therefore essential. This requires the determination of proper relieving conditions (e.g., flows, pressures ,and temperatures) so that relief valves and their associated pipework can be properly sized for the worst scenario. Designers are responsible to obviate all conceivable conditions, and good judgment must be exercized in setting the relief requirements (set pressure, for example) in all design facets during hazard and operability studies. Here, five relief conditions are considered [11]:

- gas or vapor
- steam
- liquid

- air
- fire

In addition, noise suppression, preferred location of relief valves, and pressure drop requirements for stable operation involving inlet pipe to the valve and tail pipe are reviewed.

Typical Relief Valves

There are two main types of safety relief valves. The conventional type is shown in Figure 5-3. A balanced safety relief valve shown in Figure 5-4 is designed to limit the effect of back pressure on opening pressure, closing pressure, lift capacity, and relieving capacity. The balanced valves are mainly the piston type and the bellows type.

Because of its design, a conventional relief valve encounters any back pressure accumulated in the discharge header so that the relieving pressure is affected. For example, a relief valve set at 450 psig relieving into a closed header system with 10 psig back pressure caused by loads from other units will not open until the vessel is at 460 psig. Normal practice allows only 10 percent of the set pressure "buildup" in a unit for most emergencies. Conventional relief valves, therefore, may often force the designer to increase the header size to limit back pressure. The balanced relief valve does not encounter the accumulated pressure and can relieve at design rates even when back pressure is 30–50 percent of its set pressure. Therefore, for services with high back pressure in the header, the balanced relief valve is much preferred.

Gas or Vapor Sizing

The expression used to determine the relief area for vapor discharge when the back pressure is less than the critical flow pressure is:

$$A = \frac{W}{(C)(K)(P)(K_{b})} \left[\frac{T.Z}{M_{w}}\right]^{0.5}$$
(5-46)

where C can be expressed as:

$$C = 520 \left[k \left(\frac{2}{k+1} \right)^{\frac{(k+1)}{(k-1)}} \right]^{0.5}$$
(5-47)

CONVENTIONAL SAFETY-RELIEF VALVE

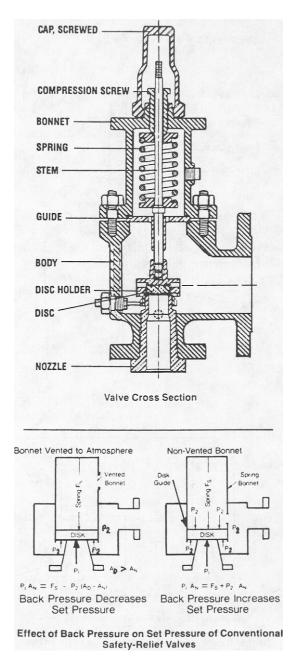
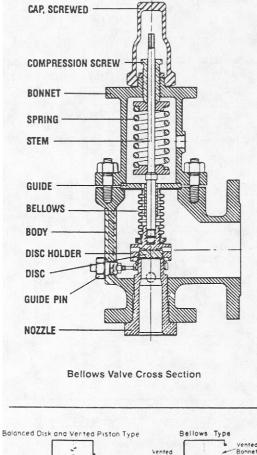
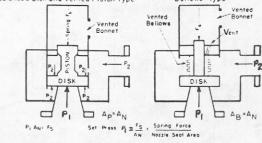


Figure 5-3. Conventional safety relief valve. (Reprinted courtesy of the American Petroleum Institute.)

BALANCED SAFETY-RELIEF VALVE





NOTE: Back pressure has very little effect on set pressure.

Effect of Back Pressure on Set Pressure of Balanced Safety-Relief Valves

Figure 5-4. Balanced safety-relief valve. (Reprinted courtesy of the American Petroleum Institute.) The values of k for some gases are given in Chapter Three. K_b is the back pressure correction factor due to back pressure and is dependent on the type of relief used. Figure 5-5 gives K_b for conventional spring reliefs and Figure 5-6 for balanced bellows reliefs, repectively.

Steam Sizing

The equation for steam service is a modification of Napier steam flow given in the ASME Boiler and Pressure Vessel Code. ASME Power Boiler Code applications permit 3 percent overpressure. The equation for steam relief is [12]:

$$A = \frac{W}{(51.5)(P_1)(K_d)(K_N)(K_{SH})}$$
(5-48)

Liquid Sizing

The requisite orifice area for a safety relief valve used in liquid service (nonviscous) requiring liquid capacity certification is:

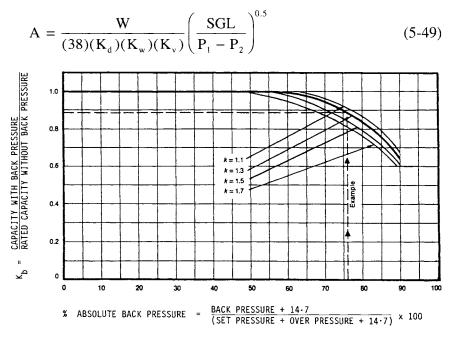


Figure 5-5. Constant back pressure sizing factor, K_p, for conventional safety relief valves in vapor or gas service. (API Recommended Practice 520, Sizing, Selection and Installation of Pressure Relieving Devices in Refineries, Part 1, 5th ed., 1990. Reprinted courtesy of the American Petroleum Institute.)

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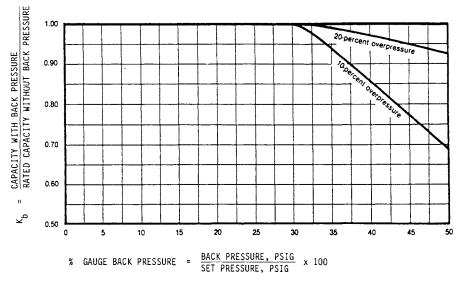


Figure 5-6. Back pressure sizing factor, K_b, for balanced-bellows pressure relief valves in vapor or gas service. (API Recommended Practice 520, Sizing, Selection and Installation of Pressure Relieving Devices in Refineries, Part 1, 5th ed., 1990. Reprinted courtesy of the American Petroleum Institute.)

 K_w is the capacity correction factor due to back pressure on balancedbellow relief valves. If the back pressure is atmospheric $K_w = 1$. This correction factor is given is Figure 5-7. When sizing a relief valve for viscous liquid service, the orifice area is first calculated for nonviscous service to obtain a preliminary discharge area. The next standard orifice size is used to calculate the Reynolds number. The K_v factor is then calculated and applied to correct the preliminary required discharge area. The Reynolds number is expressed as [10]:

$$Re = \frac{Q(2800)(SGL)}{\mu(ADES)^{0.5}}$$
(5-50)

where

$$K_v = -2.38878 + 1.2502(\ln \text{Re}) + 0.1751(\ln \text{Re})^2 + 0.01087(\ln \text{Re})^3 - 0.00025(\ln \text{Re})^4$$
(5-51)

Air Sizing

The required orifice area for a relief valve sizing for air at constant back pressure is:

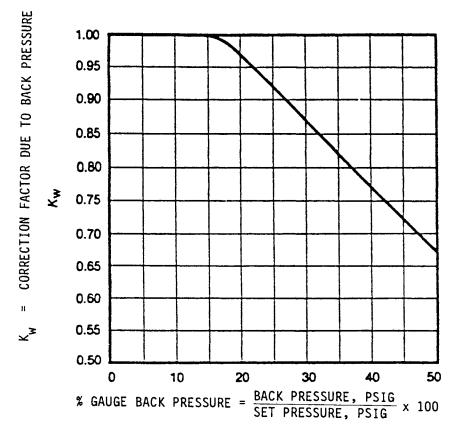


Figure 5-7. Capacity correction factor, K_w, due to back pressure on balanced-bellows pressure relief valves in liquid service. (API Recommended Practice 520, Sizing, Selection and Installation of Pressure Relieving Devices in Refineries, Part 1, 5th ed., 1990. Reprinted courtesy of the American Petroleum Institute.)

$$A = \frac{(V_a)(T)^{0.5}}{(418)(K_d)(P)(K_b)}$$
(5-52)

Fire Sizing

In the case of a fire, there is a pressure buildup and the required relief capacity can be determined provided that the heat absorbed can be estimated. API 520 gives recommendations for heat of absorption during vapor relief [12]. Not all the causes will happen simultaneously, but the pressure relief or safety relief should be sized for conditions that require the greatest relief.

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Inconsistencies for sizing relief valves for fire emergencies arise among published codes. This is because of varying interpretations of the heat flux caused by fire exposure. These variations are based on the area in which the heat flux applies and the protection factors used to characterize the installations. Crozier [13] has summarized these variations and evaluated the pertinent correlations. Here, the API 520 correlation is used because it contains an implied protection factor of 0.5 for good drainage.

Liquid in liquid-filled vessels exposed to direct or radiated heat from a fire will vaporize. The heat required to accomplish this will limit the shell temperature of the tank to only a slight rise. The amount of liquid vaporized depends on the rate of heat input from the fire and the relieving temperature. This can be determined by calculating the heat input to the wetted surface of the vessel and dividing this by the latent heat of vaporization.

$$Q_r = 21,000(FI)(A_w)^{0.82}$$
(5-53)

and

$$W = \frac{Q_r}{\lambda}$$
(5-54)

The vapor to be relieved is the vapor in equilibrium with the liquid (that is, saturated vapor at the conditions existing when the valve is relieving at the rated capacity). Table 5-3 shows environmental factors for bare and insulated vessels.

For a multicomponent mixture, λ_{mix} can be determined from

$$\lambda_{mix} = \sum_{i=1}^{n} \frac{\lambda_i y_i M_i}{M}$$
(5-55)

To determine y_i , it is necessary to know the equilibrium liquid-phase mole fraction, x_i , at the bubble point corresponding to the accumulated relieving pressure.

The wetted surface area depends on the diameter and length of the vessel. For working storage tanks, the wetted surface is obtained on the average inventory. Surge drums usually operate about half full; thus the wetted surface will be calculated at 50 percent of the total vessel surface.

Type of Equipment	Factor
Bare vessel	1.0
Insulated vessel (These arbitrary insulation conductance values are in Btu/hrft ² °F	
4	0.3
2	0.15
1	0.075
0.67	0.05
0.5	0.0376
0.4	0.03
0.33	0.026
Water-application facilities on bare vessel	1.0
Depressurizing and emptying facilities	1.0
For approved water spray with approved insulation	0.15
For drainage	0.5
For approved water spray	0.3

Table 5-3 Environmental Factor

Effect of Back Pressure

The maximum allowable back pressure (gauge) for a conventional safety relief valve is 10% of the set pressure (gauge). If the maximum allowable back pressure is exceeded during any relief situation, the size of the relief lines or headers should be increased until the back pressures are sufficiently reduced.

Noise Suppression

Expansion of gas or vapor usually gives rise to an unacceptable noise level. However, this is not a problem with liquid relief. There are two sources of noise:

- 1. Noise due to turbulence at the point where the gas mixes with the atmosphere. Suppression of this type of noise involves the fitting of a silencer.
- 2. Noise due to pressure letdown (that is, relief valve) when sonic or near sonic velocities are generated. This is transmitted through the wall of pipe and, if the discharge is short, it is transmitted through the open end. Noise from this can be reduced by lengthening the discharged pipe and by insulation. Suppression of the noise can also be overcome by fitting a silencer. Such silencers are often installed in an exposed environment and are likely to accumulate dirt, vapor, or products from minor leakage of safety devices.

Effect of Pressure Drop (ΔP) on Pipework

The pipework should be designed with the following considerations:

- The frictional ΔP through the inlet pipework, inlet fittings, and valves at the maximum possible relief rate should not exceed 3% of the set gauge pressure (psig). The calculation of ΔP should include the entrance loss of the inlet pipework. The discharge pipework must be such that ΔP is less than 10% of the relieving pressure (that is, set pressure plus overpressure in gauge). The discharge pipework should withstand the internal pressure and high velocities, the reaction forces, and the sudden transient loading that occur when a relief device suddenly comes into operation.
- The inlet pipework should be of a size at least equal to the relief valve, with its length minimized to reduce ΔP . It should also be equal to the bending moments resulting from the forces that develop when the valve or disc is discharging at full capacity.
- A relief device should be installed vertically and preferably on a nozzle at the top of the equipment (for example, vessel) or on a tee connected to a pipeline. When discharging a gas or vapor, the fluid reaches sonic velocity when passing through a relief device. Thus, the gas flow rate or ΔP can be determined by a method as described in Chapter Three.

For liquid discharges, the design of the pipework may be complex. If the liquid is subcooled, the discharge pipework is sized by standard liquid pressure formulae. If the liquid in the equipment is at or near its saturation pressure, the relief device and discharge pipework should be sized for two-phase flow. As the liquid flows through the relief valve, the ΔP will cause it to flash. Therefore, the size of the valve and discharge pipework must be increased to accommodate the larger volume. Further flashing in the downstream piping will increase the gas-liquid mixture through the two-phase regimes. If this is not arrested, slug flow will develop along the pipeline resulting in the mechanical damage to the pipework.

Discharge Reactive Forces

The total stress imposed on a safety valve or its piping is a result of the sum of the following:

- internal pressure
- dead weight of piping
- thermal expansion or contraction of either the discharge line
- the equipment upon which the valve is mounted
- the bending moment caused by the reaction thrust at the discharge

The discharge reactive force is based on the assumption that critical flow of the gas or vapor is obtained at the outlet of the relief device and the discharge is horizontal to the atmosphere. For any gas or vapor, the reactive force can be expressed as:

$$F = \frac{W \left[\frac{kT}{(k+1)M_{w}}\right]^{0.5}}{366}$$
(5-56)

Relief Valve Location

The location of a relief device often influences its size and material of construction. Relief devices must be sited so that the risk of blockage by debris is minimized. It is poor practice to mount relief valves at the end of long horizontal pipe through which there is no flow. The possibility of liquid phase entering a vapor relief system must also be avoided. Locating a relief device upstream of equipment that has ΔP due to flow allows the relief valve to be smaller because increased ΔP is available. Reduced costs of a relief system may be obtained by installing relief valves in a more conducive position with regard to temperature and corrosion. Figure 5-8 shows alternative positions. Position 1 is preferred when the relief valve is tied into a closed relief system because of ease of maintenance. It assures a relatively short line to the relief header and

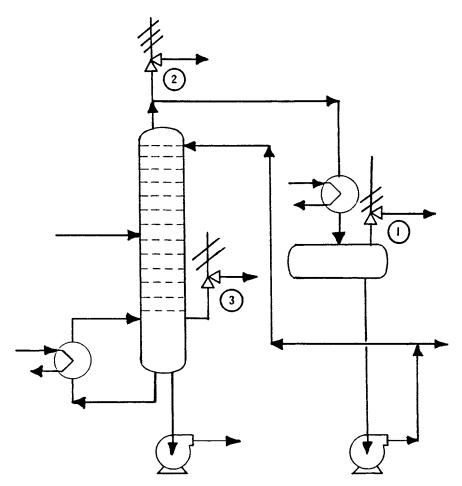


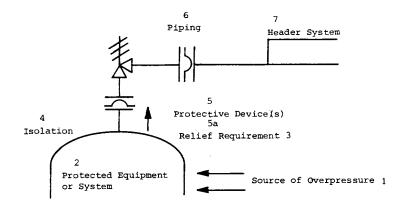
Figure 5-8. Alternative positions for a relief valve.

lower temperature for relief when air fan coolers are used. Position 2 is preferred for relief to atmosphere. Position 3 is used on low pressure towers designed for the same top and bottom pressure. Figure 5-9 illustrates key areas for design verification of relief system. Manufacturers' orifice areas and corresponding valve sizes are shown in Table 5-4.

Nomenclature

A = valve discharge area, sq in.

- A_w = wetted surface area of the vessel, sq ft
- ADES = next largest valve orifice, sq. in, from the manufacturer's standard orifice area



	KEY AREA	BASIC CONSIDERATIONS
1.	Sources of Overpressure	Which have been considered?
2.	Equipment or System protected by Relief stream	 (i) All equipment is identified. (ii) Design press. and temp. limits cannot be exceeded Design codes for protected equipment and devices are compatible.
з.	Require Relief Rate	What is it?
4.	Isolation	Is any fitted? If so, how is continuity assured?
5.	Protective Device(s)	Under or oversized? Suitability of device(s).
	5a. Devices in Series	Correct ΔP maintained across devices. Interspace monitored/alarmed
6.	Piping	Correctly sized. Pressure drop O.K. Supports O.K for reaction loads. Suitable for fluid. Drainage/blockage.
7.	Headers or Common Vents	Is there one? Sized for all streams. Back pressure O.K. Damage to/from other streams.

Figure 5-9. Pressure relief design verification.

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Orifice Designation	Orifice Area (sq. in.)	Valve Size (in.)
D	0.110	1–2
E	0.196	1-2
F	0.307	1.5-2
G	0.503	1.5-2.5
Н	0.785	1.5-3 or 2-3
J	1.287	2–3 or 2.5–4
K	1.838	3–4
L	2.853	3–4 or 4–6
Μ	3.600	4-6
Ν	4.340	46
Р	6.379	4-6
Q	11.050	6–8
R	16.000	6–8 or 6–10
Т	26.000	8-10

Table 5-4Orifice Sizes for Relief Valves

- C = coefficient determined by ratio of specific heats of the gas or vapor at standard conditions
- F = reaction force at the point of discharge to the atmosphere, lb
- FI = factor for insulation (for non-insulated vessel FI is 1.0)
- K = effective coefficient factor due to backpressure (0.975 for vapor flow)
- K_b = capacity correction factor due to backpressure is 1.0, when backpressure is 55% of absolute relieving pressure
- K_d = coefficient of discharge (0.953)
- $K_N = correction factor for Napier equation, K_N$, is 1.0, where $P \le 1515 psia$
- K_p = overpressure correction factor for direct spring-operated valves only, K_p is 1.0
- K_{SH} = correction factor due to amount of superheat in the steam. Values range from 0.88 (maximum superheat) to 1.0 for saturated steam.
 - $K_v =$ viscosity correction factor
 - K_w = backpressure correction factor for balanced-bellows springoperated valves only (K_w is 1.0)
 - k = ratio of specific heat capacities (C_p/C_v)

- M = molecular weight
- M_w = molecular weight of vapor (or process fluid)
 - P = upstream relieving pressure, psia
 - P_{b} = total backpressure, psig
 - P_i = set pressure at which relief valve begins to open, psig
 - Q = flow rate, in U.S. gallons per min
 - Q_r = rate of heat input due to fire, Btu/hr
- SGL = specific gravity of liquid at flowing temperature referred to water as 1.0 at 70° F
 - T = absolute temperature of inlet vapor $^{\circ}R(^{\circ}F + 460)$
 - V_a = required air capacity, standard ft³/min
 - W =flow of any gas or vapor, lb/hr
 - Z = compressibility factor
 - λ = latent heat of vaporization at the boiling point of the liquid, Btu/ib
 - μ = absolute viscosity at the flowing temperature, cP

TWO-PHASE FLOW RELIEF SIZING FOR RUNAWAY REACTION

Introduction

Many methods have been used to size relief systems: area/volume scaling, mathematical modeling using reaction parameters and flow theory, and empirical methods by the Factory Insurance Association (FIA). The Design Institute for Emergency Relief Systems (DIERS) of the AIChE has carried studies of sizing reactors undergoing runaway reactions. Intricate laboratory equipment and analysis have resulted in better vent sizes.

A selection of relief venting as the basis of safe operation is based upon the following considerations [14]:

- compatibility of relief venting with the design and operation of the plant/process
- identifying the worst scenarios
- type of reaction
- means of measuring the reaction parameters during the runaway reaction
- relief sizing procedure
- design of the relief system including discharge ducting and safe discharge area

Runaway Reactions

A runaway reaction occurs when an exothermic system becomes uncontrollable. The reaction leads to a rapid increase in the temperature and pressure, which if not relieved can rupture the containing vessel. A runaway reaction happens because the rate of reaction and therefore the rate of heat generation increases exponentially with temperature. Alternatively, the rate of cooling increases only linearly with temperature. Once the rate of heat generation exceeds available cooling, the rate of increase in temperature becomes progressively faster. Runaway reactions nearly always result in two-phase flow reliefs. Runaway reactions are generally classified into three systems.

Vapor Systems

Boiling is attained before potential gaseous decomposition, that is, the heat of reaction is removed by the latent heat of vaporization. The reaction is tempered, and the total pressure in the reactor is equal to the vapor pressure. The principal parameter determining the vent size is the rate of the temperature rise at the relief set pressure.

Gassy Systems

Gaseous decomposition reaction occurs without tempering. The total pressure in the reactor is equal to the gas pressure. The principal parameter determining the vent size is the maximum rate of pressure rise.

Hybrid Systems

Gaseous decomposition reaction occurs before boiling. The reaction is still tempered by vapor stripping. The total pressure in the reactor is the summation of the gas partial pressure and the vapor pressure. The principal parameters determining the vent size are the rates of temperature and pressure rise corresponding to the tempering condition. A tempered reactor contains a volatile fluid that vaporizes or flashes during the relieving process. This vaporization removes energy via the heat of vaporization and tempers the rate of temperature rise due to the exothermic reaction.

Adiabatic Calorimetry

Two-phase flow calculations are complex when conditions change rapidly as in runaway reactions. Because of this complexity, several experimental and analytical techniques have been developed over the past decade to obtain the relevant data for the relief vent calculations. The data required for the runaway calculations are determined with laboratory tools such as the Differential Scanning Calorimetry (DSC), the accelerating rate calorimeter (ARC), the vent sizing package (VSP), and recently, the laboratory calorimetric Reactive System Screening Tool (RSST). The RSST determines the type of reaction and principal parameters required for sizing reactor vents for vapor, hybrid, and gassy systems. Figures 5-10A and 5-10B show the principal features of the RSST. The principal data for relief sizing calculations are shown in Figures 5-11A, 5-11B, 5-11C, and 5-11D, the temperature rate at the set *(text continued on page 366)*

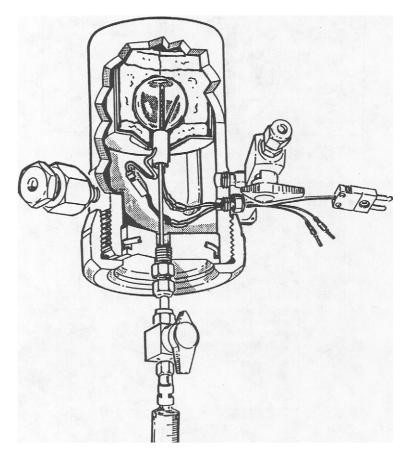


Figure 5-10A. The RSST vessel.

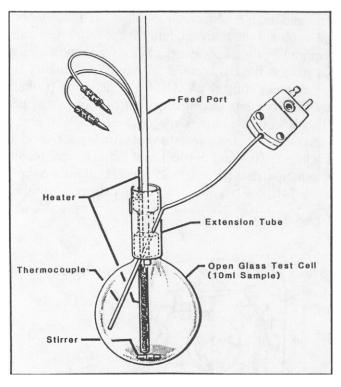


Figure 5-10B. The test cell and associated equipment.

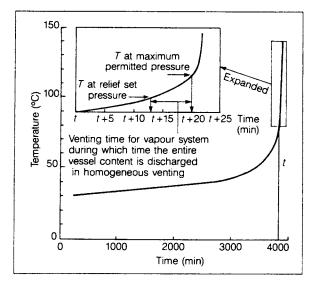


Figure 5-11A. Temperature against time for an abiabatic system.

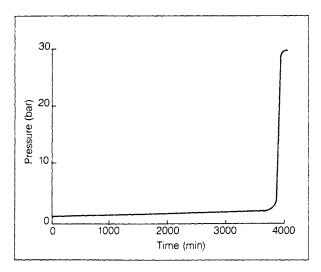


Figure 5-11B. Pressure against time for an abiabatic system.

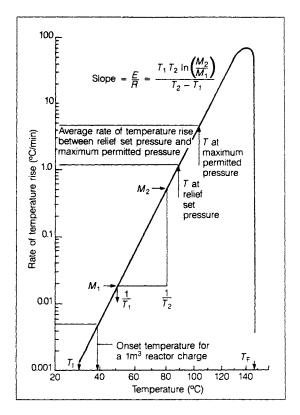


Figure 5-11C. Rate of temperature rise against temperature for an abiabatic system.

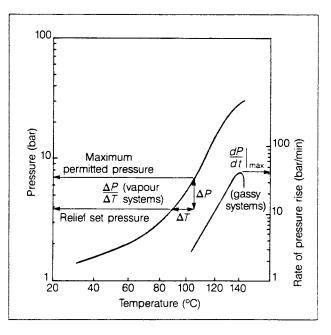


Figure 5-11D. Pressure and rate of pressure rise against temperature for an abiabatic system.

(text continued from page 361)

pressure and the temperature increase, corresponding to the overpressure. In addition, the heat of reactions can be determined from the temperature vs. time data of the heat capacities if the monomers and products are available. Table 5-5 lists formulae for computing the area of the three systems.

Simplified Nomograph Method

Boyle [15] and Huff [16] first accounted for two-phase flow with relief system design for runaway chemical reactions. A computer simulation approach to vent sizing involves extensive thermokinetic and thermophysical characterization of the reaction system. Fisher [17] has provided an excellent review of emergency relief system design involving runaway reactions in reactors and vessels. Fauske [18] has developed a simplified chart to the two-phase calculation. He expressed the relief area as:

$$A_{I} = \frac{V\rho_{r}}{G\Delta t_{v}}$$
(5-57)

Vapor System Gassy System Hybrid System $A = 5.6 \times 10^{-6} \left(\frac{1}{F}\right) \left(\frac{m_0}{m_t}\right) \left(\frac{\frac{dP}{dt}}{P_s^{1.5}}\right)$ $A = 1.5 \times 10^{-5} \left(\frac{m_0 \frac{d\Gamma}{dt}}{F.P_s} \right)$ $A = 3 \times 10^{-6} \left(\frac{1}{F}\right) \left(\frac{m_0}{m_t}\right) \left(\frac{\frac{dP}{dt}}{P_{MAP}^{1.5}}\right)$ $d = \left(\frac{4A}{\pi}\right)^{0.5}$ $\mathbf{d} = \left(\frac{4\,\mathrm{A}}{\pi}\right)^{0.5}$ $d = \left(\frac{4A}{\pi}\right)^{0.5}$ L/D=0, F=1.00 L/D=0F=1.00 L/D=0, F=1.00 L/D=50, F=0.85 L/D=50, F=0.70 L/D=50, F=0.70 L/D=100, F=0.75 L/D=100. F=0.60 L/D=100, F=0.60 L/D=200. F=0.65 L/D=200. F=0.45 L/D=200, F=0.45 L/D=400, F=0.50 L/D=400, F=0.33 L/D=400, F=0.33

 Table 5-5

 Vent Areas and Diameters of the Three Systems

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The mass flux can either be expressed as:

$$G \cong 0.9F_1 \frac{\Delta P}{\Delta T} \left(\frac{g_c T_s}{C_p}\right)^{0.5}$$
(5-58)

or

$$G \cong 0.9F_{I} \frac{\Delta H_{v}}{v_{fg}} \left(\frac{g_{c}}{C_{p}T_{s}}\right)^{0.5}$$
(5-59)

The venting time is given by

$$\Delta t_{v} \cong \frac{(\Delta T)(C_{p})}{q_{s}}$$
(5-60)

Combining Equations 5-58, 5-59, and 5-60, the area, A, is given by

$$A = \frac{V\rho q_s}{\Delta P} \left(g_c C_p T_s \right)^{-0.5}$$
(5-61)

Equation 5-61 gives a conservative estimate of the vent area and the simple design method represents overpressure (ΔP) between 10 and 30 percent. For a 20 percent absolute overpressure, a liquid heat capacity of 2510 J/kg.K for most organics, and considering that a saturated water relationship exists, the vent size area per 1,000 kg of reactants is:

$$A = \left(\frac{m^2}{1,000 \text{ kg}}\right) = \frac{0.00208 \left(\frac{dT}{dt}\right)}{P_s} \frac{\text{°C/min.}}{\text{bar}}$$
(5-62)

Figure 5-12 shows a nomograph of determining the vent size. The vent area is calculated from the heating rate, the set pressure, and the mass of reactants. The nomograph is used for obtaining quick vent sizes and checking the results of the more rigorous computation. Crowl and Louvar [19] have expressed that the nomograph data of Figure 5-12 applies for a discharge coefficient of F = 0.5, representing a discharge L/D of 400.0. However, use of the nomograph at other discharge pipe lengths and different F requires a suitable conversion.

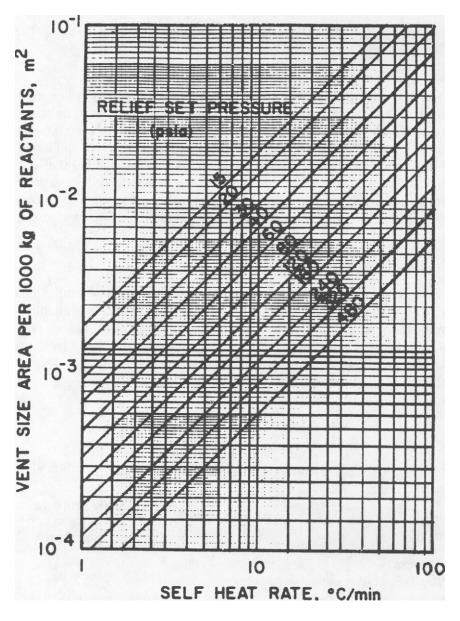


Figure 5-12. A vent sizing nomogram for tempered (high vapor pressure) runaway chemical reactions. (Courtesy of the American Institute of Chemical Engineers.)

Vent Sizing Methods

Vents are usually sized on the assumption that the vent flow is:

- all vapor or gas
- all liquid
- a two-phase mixture of liquid and vapor or gas

The first two cases represent the smallest and largest vent sizes required for a given rate at increased pressure. Between these cases, there is a two-phase mixture of vapor and liquid. It is assumed that the mixture is homogeneous, that is, that no slip occurs between the vapor and liquid. Furthermore, the ratio of vapor to liquid determines whether the venting is closer to the all vapor or all liquid case. As most relief situations involve a liquid fraction of over 80 percent, the idea of homogeneous venting is more to all liquid than all vapor. Table 5-6 shows vent area for different flow regimes.

Vapor Pressure Systems

These systems are called "tempering" (that is, to prevent temperature rise after venting) systems because sufficient latent heat is available to remove the heat of reaction and to temper the reaction at the set pressure. The vent requirements for such systems are estimated from the Leung's Method [20,21].

Type of flow	Required vent area as a multiple of all vapor vent area
All vapor	1
Two-phase: churn turbulent	2–5
Bubbly	7
Homogeneous	8
All liquid	10

Table 5-6Vent Areas for Different Flow Regimes

$$A = \frac{M_0 q}{G \left[\left(\frac{V}{M_0} T_s \frac{dP}{dT} \right)^{0.5} + (C\Delta T)^{0.5} \right]^2}$$
(5-63)

Alternatively, the vent area can be expressed as:

$$A = \frac{M_0 q}{G \left[\left(\frac{V}{M_0} \frac{h_{fg}}{v_{fg}} \right)^{0.5} + (C\Delta T)^{0.5} \right]^2}$$
(5-64)

The vapor pressure systems obey the Antoine relationships:

$$\ln P = A + \frac{B}{T}$$
(5-65)

differentiating Equation 5-65 yields,

$$\frac{1}{P}\frac{\mathrm{d}P}{\mathrm{d}T} = -\frac{B}{T^2} \tag{5-66}$$

$$\frac{\mathrm{dP}}{\mathrm{dT}} = -\frac{\mathrm{B}}{\mathrm{T}^2} \,\mathrm{P} \tag{5-67}$$

Fauske's Method

Fauske represented a nomograph for tempered reactions as shown in Figure 5-12. This accounts for turbulent flashing flow and requires information about the rate of temperature rise at the relief set pressure. This approach also accounts for vapor disengagement and frictional effects including laminar and turbulent flow conditions. For turbulent flow, the vent area is

$$A = \frac{1}{2} \frac{M_0 \left(\frac{dT}{dt}\right)_s (\alpha_D - \alpha_0)}{F_2 \left(\frac{T_s}{C_s}\right)^{0.5} \Delta P(1 - \alpha_0)}$$
(5-68)

Figure 5-13 shows a sketch of temperature profile for high vapor pressure systems.

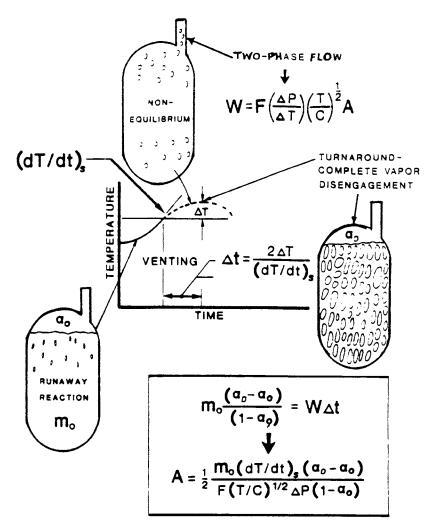


Figure 5-13. Vent sizing model for high vapor pressure systems. Due to non-equilibrium effects, turnaround in temperature is assumed to coincide with the onset of complete vapor disengagement.

Gassy Systems

The major method of vent sizing for gassy system is two-phase venting to keep the pressure constant. This method was employed before DIERS with an appropriate safety factor [22]. The vent area is expressed by:

$$A = \frac{Q_g(1-\alpha)\rho_f}{G_1} = \frac{Q_g M}{GV}$$
(5-69)

Unlike systems with vapor present, gassy systems do not have appreciable latent heat to temper the reaction. The system pressure increases as the rate of gas generation with temperature increases, until it reaches the maximum value. The vent area could be underestimated if sizing is dependent on the rate of gas generated at the set pressure. Therefore, it is more plausible to size the vent area on the maximum rate of gas generation. Homogeneous two-phase venting is assumed even if the discharge of liquid during venting could reduce the rate of gas generation even further. The vent area is defined by

$$A = 3.6 \times 10^{-3} Q_{g} \left(\frac{M_{0}}{VP_{m}}\right)^{0.5}$$
(5-70)

The maximum rate of gas generation during a runaway reaction is proportional to the maximum value of dP/dt and can be calculated from

$$Q_{gl} = \frac{M_0}{M_t} \left(\frac{V_t}{P_m} \frac{dP}{dt} \right)$$
(5-71)

Homogeneous Two-Phase Venting Until Disengagement

Imperial Chemical Industry (ICI) [22] developed a method for sizing a relief system that accounts for vapor-liquid disengagement. They proposed that homogeneous two-phase venting occurs that increases to the point of disengagement. Furthermore, they based their derivations upon the following assumptions:

- vapor phase sensible heat terms may be neglected.
- vapor phase mass is negligible.
- heat evolution rate per unit mass of reactants is constant (or average value can be used).
- mass vent area per unit area is approximately constant.
- physical properties can be approximated by average values.

The vent area is given by the following equation.

$$A = \frac{q_1 V(\alpha - \alpha_0)}{G_1 v_f \left\{ \frac{h_{fg} v_f(\alpha - \alpha_0)}{v_{fg}(1 - \alpha_0)(1 - \alpha)} + C\Delta T_1 \right\}}$$
(5-72)

Equation 5-72 is valid if disengagement occurs before the pressure would have turned over during homogeneous venting; otherwise, Equation 5-72 gives an unsafe (too small) vent size. Therefore, it is necessary to verify that

$$q > \frac{G_1 A h_{fg} v_f^2}{V v_{fg} (1 - \alpha)^2}$$
(5-73)

is satisfied at the point of disengagement.

Two-Phase Flow Through An Orifice

Sizing formulae for flashing two-phase flow through relief devices were obtained through DIERS, which is based upon Fauske's equilibrium rate model (ERM) and assumes frozen flow (non flashing) from a stagnant vessel to the relief device throat. This is followed by flashing to equilibrium in the throat. The orifice area, A, is given by,

$$\mathbf{A} = \frac{\mathbf{W}}{\mathbf{C}_{\mathrm{D}}} \left\{ \left(\frac{\mathbf{x} \mathbf{V}_{\mathrm{G}}}{\mathbf{k} \mathbf{P}_{\mathrm{I}}} \right) + \left[\frac{\left(\mathbf{V}_{\mathrm{G}} - \mathbf{V}_{\mathrm{L}} \right)^{2} \mathbf{C}_{\mathrm{L}} \mathbf{T}_{\mathrm{I}}}{\lambda^{2}} \right] \right\}^{0.5}$$
(5-74)

The theoretical rate can be expressed as:

$$W = A_f (2\Delta P.\rho)^{0.5}$$
(5-75)

For a simple sharp-edged orifice, the value of C_D , the coefficient of discharge, is well established (about 0.6). For safety relief valves, its value depends upon the shape of the nozzle and other design features. In addition, the value C_D varies with the conditions at the orifice. For saturated liquid at the inlet, Equation 5-74 simplifies to the following expression.

$$A = \frac{W(V_{g} - V_{L})(C_{L}T_{1})^{0.5}}{C_{p}\lambda} = \frac{W}{C_{p}\left(\frac{dP}{dT}\right)\left(\frac{T_{1}}{C_{L}}\right)^{0.5}}$$
(5-76)

Equations 5-72 to 5-76 are based upon the following assumptions:

- vapor phase behaves as an ideal gas.
- liquid phase is an incompressible fluid.
- turbulent Newtonian flow is present.

• critical flow. This is usually the case because the critical pressure ratios of flashing liquid approach the value of 1.0.

Duxbury and Wilday [22] recommend that a safety factor of at least 2.0 be used. In certain cases, lower safety factors may be employed. However, the designer should consult the appropriate process safety section in the engineering department for advice.

Discharge System

Design of the Vent Pipe. The nature of the discharging fluid is necessary in determining the relief areas. DIERS and ICI techniques can analyze systems that exhibit "natural" surface active foaming and those that do not. DIERS further found that small quantities of impurities can affect the flow regime in the reactor. In addition, a variation in impurity level could arise by changing the supplier of a particular raw material. Therefore, care is needed in sizing emergency relief on homogeneous vessel behavior, that is, two-phase flow.

In certain instances, pressure relief during a runaway reaction can result in a three-phase discharge if solids are suspended in the reaction liquors. Solids can also be entrained by turbulence caused by boiling or gassing in the bulk of the liquid. Caution is required in sizing this type of relief system, especially where there is a significantly static head of fluid in the discharge pipe. Another aspect in the design of the relief system includes the possible blockage in the vent line. This could arise from the process material being solidified in cooler sections of the reactor. It is important to consider all discharge regimes when designing the discharge pipe work.

Safe Discharge

Reactors or storage vessels are fitted with overpressure protection vent directly to roof level. Such devices (e.g., relief valves) protect only against common process maloperations and not runaway reactions. The quantity of material ejected and the rate of discharge are low, resulting in good dispersion. The rise of bursting disc can result in large quantities of materials (95% of the reactor contents) being discharged for foaming systems.

The discharge of copious quantities of chemicals directly to atmosphere can give rise to secondary hazards, especially if the materials are toxic and can form flammable atmosphere in air (e.g., vapor or mist). In such cases, the provision of a knock-out device (scrubber, dump tank) of adequate size to contain the aerated/foaming fluid will be required.

Conditions of Use

- If Fauske's method yields a significantly different vent size, then the calculation should be reviewed.
- The answer obtained from the Leung's method should not be significantly smaller than that from Fauske's method.
- ICI recommends a safety factor of 1–2 on flow or area. The safety factor associated with the inaccuracies of the flow calculation will depend on the method used, the phase nature of the flow, and the pipe friction. For two-phase flow, use a safety factor of two to account for friction and static head.
- Choose the smaller of the vent size from the two methods.

A systematic evaluation of venting requires information on:

- the reaction—vapor, gassy, or hybrid.
- flow regime—foamy or nonfoamy.
- vent flow—laminar or turbulent.
- vent sizing parameters, such as dT/dt, $\Delta P/\Delta T$, ΔT .

It is important to ensure that all factors (e.g., long vent lines) are accounted for, independent of the methods used. Designers should ascertain that a valid method is chosen rather than the most convenient or the best one. For ease of use, the Leung's method for vapor pressure systems are rapid and easy. Different methods should give vent sizes with a factor of two. Nomographs give adequate vent sizes for long lines to L/D of 400, but sizes are divisible by two for nozzles. Table 5-6A lists the applicability of both Leung and Fauske's methods.

Nomenclature

- A = vent area, m^2
- $A_1 = relief vent area$
- $A_f = flow area, m^2$
- \dot{C} = specific heat of the material, J/Kg.K
- $C_{\rm D}$ = coefficient of discharge ($C_{\rm D}$ = 0.6)
- $\overline{C_L}$ = average liquid specific heat capacity, J/Kg.K
- C_p = specific heat at constant pressure, J/Kg.K
- d = vent diameter, mm
- F = flow reduction factor
- F_1 = correction factor, and for a length of pipe zero, F_1 =1.0. As the pipe length increases, the value of F_1 decreases.

Leung's Method	Fauske's Method
1. The overpressures are limited in the range 0–50% of absolute set pressure, i.e., $0 \le \Delta P \le 0.5 P_s$	The overpressures are limited in the range of 10–30% of absolute set pressure, i.e., $0 \le \Delta P \le 0.3P_s$. It assumes that flow is turbulent and that the vapor behaves as an ideal gas.
2. The method includes the mass unit vent flow capacity per unit area, G. This allows the user to select any applicable vent capacity calculation method.	The method incorporates the equilibrium rate model for vent flow capacity when friction is negligible. In addition, a correction factor is used for longer vent lines of constant diameter and with negligible static head change.
3. The heat evolution rate per unit mass, the vent capacity per unit area, physical properties (e.g., latent heat of fluid, specific heat and vapor/liquid specific volumes) are constant.	It allows for total vapor-liquid disengagement of fluids that are not "natural" surface active foamers.
4. A hand calculation method that can be used to take account of two- phase relief when the materials in the vessel are "natural" surface active foamers.	To account for disengagement, the vessel void fraction at disengage- ment should be evaluated (i.e., the point at which the vent flow would cease to be two-phase and start to be vapor only).
5. The methods use drift-flux level- swell calculation models to take account of more vapor being in the inlet stream to relief device than the average for the vessel.	The method is potentially unsafe if early disengagement (before the pressure would otherwise have turned over during two-phase venting) occurs.
6. Methods are given for the bubbly and churn-turbulent flow regimes within the vessel.	The criterion for safe use to take account of disengagement is $q < \frac{G A h_{fg} v_f^2}{V v_{fg} (1 - \alpha_D)^2}.$ However, when disengagement is not accounted for, then $\alpha_D = 1.0$, and the above equation is not required.

Table 5-6A Applicability of Recommended Vent Sizing Methods

 F_2 = flow reduction factor for turbulent flow

 \overline{G} = mass flux through the relief

 G_1 = mass vent capacity per unit area, kg/m²s

 g_c = dimensional gravitational constant, 1.0 kg m/N s²

 h_{fg} = latent heat, J/kg

 ΔH_v = heat of vaporization of the fluid, J/kg

k = isentropic coefficient

M = mass of liquid in vessel, kg

 $M_o = initial mass of liquid in vessel, kg$

 $m_t = mass of the sample in the RSST, kg$

 P_m = maximum allowable pressure during venting, bar

 P_{MAP} = maximum allowable vessel pressure, psia

 P_s = relief set pressure, psia

 P_1 = pressure in the upstream vessel, N/m²abs

 ΔP = overpressure, bar

 ΔP_1 = difference between the (reduced) set pressure and the maximum pressure allowable, N/m²

 ΔP_2 = pressure difference, N/m²

dP/dt = maximum rate of pressure rise in the test vessel, bar/s

dP/dt = measured maximum rate of pressure rise in the RSTT, lb/in² min

 Q_g = volumetric gas generation rate of temperature and pressure in reactor during relief, m³/s

 Q_{gl} = maximum rate of gas generation, m³/s

 $q_1 =$ self-heat rate, W/kg

q = rate of heat production (W/kg) at the relief set pressure, or an average value between the relief set pressure and the maximum permitted pressure if this is more than 120% of the relief set pressure

T = temperature corresponding to relief actuation, K

 T_s = absolute saturation temperature of the fluid at the set pressure, K

 ΔT = temperature rise corresponding to the overpressure, K

 ΔT_1 = temperature difference between that at the maximum pressure allowable, and that at the redefined set pressure, k.

 $\Delta t_v = venting time$

 $(dT/dt)_s$ = rate of temperature rise corresponding to the relief set pressure, K/s

 $V = reactor volume, m^3$

= total vessel volume, m³

 v_{fg} = difference between vapor and liquid specific volume, m³/kg

 $v_f =$ liquid specific volume, m³/kg.

 V_G = specific volume of gas, m³/kg.

 V_L = specific volume of liquid, m³/kg.

W = required relief rate, kg/s

x = mass fraction of vapor at inlet

 α = void fraction in vessel

- $\alpha_{\rm D}$ = vessel void fraction corresponding to complete vapor disengagement
- α_{0} = initial free board volume
- $\rho = \text{density}, \text{kg/m}^3$
- $\rho_f =$ liquid density, kg/m³

 ρ_r = density of the reactants, kg/m³

PROBLEMS AND SOLUTIONS

Problem 5.1 Orifice Sizing for Liquid Flow

Calculate the size of an orifice for the bore of a flange-tapped orifice required to measure 12.84 ft³/min of gasoline in a 2-inch schedule 80 pipe.

Design data:

Internal pipe diameter (I.D.)	= 1.939 inches
Normal differential pressure	= 64 in H2O
Flowing temperature	= 80.5°F
Density of gasoline	$= 42.26 \text{ lb/ft}^3$
Viscosity of gasoline	= 0.6 cP
Assuming the coefficient of discharge C _D	= 0.62
F _a	= 1.0

Orifice Sizing for Gas Flow

Determine the size of an orifice to give a full-scale reading of 20 in. H_2O for 0.25 lb/s flow of gas through a 4-inch (I.D = 4.026) schedule 40 pipe. The upstream gas density is 0.075 lb/ft³, viscosity of the gas is 0.0219 cP, the specific heat ratio, k, is 1.4, and the upstream pressure is 14.7 psi.

Solutions

The computer program **PROG51** calculates the orifice size for liquid and gas services. Tables 5-7 and 5-8 show the input data and results of the liquid and gas orifice calculations. For the liquid service of gasoline,

Table 5-7 Orifice Sizing for Liquid Flow: Program Inputs and Outputs

LIQUID INCH-H20 1.939 64.0 42.26 12.84 0.6 ORIFICE SIZING FOR LIQUID FLOW TYPE OF FLOW: TYPE OF PRESSURE DROP: INCH-H20 INTERNAL PIPE DIAMETER, inch: 0RIFICE PRESSURE DROP, inch-H20: 0RIFICE PRESSURE DROP, psi: 2.308 LIQUID DENSITY, lb/ft^3: 42.260
INCH-H20 1.939 64.0 42.26 12.84 0.6 ORIFICE SIZING FOR LIQUID FLOW ************************************
1.939 64.0 42.26 12.84 0.6 ORIFICE SIZING FOR LIQUID FLOW TYPE OF FLOW: TYPE OF PRESSURE DROP: INTERNAL PIPE DIAMETER, inch: INTERNAL PIPE DIAMETER,
42.26 12.84 0.6 ORIFICE SIZING FOR LIQUID FLOW TYPE OF FLOW: TYPE OF PRESSURE DROP: INCH-H20 INTERNAL PIPE DIAMETER, inch: ORIFICE PRESSURE DROP, inch-H20: ORIFICE PRESSURE DROP, psi: LIQUID DENSITY, lb/ft^3: 42.260
ORIFICE SIZING FOR LIQUID FLOW TYPE OF FLOW: LIQUID TYPE OF PRESSURE DROP: INCH-H2O INTERNAL PIPE DIAMETER, inch: 1.939 ORIFICE PRESSURE DROP, inch-H2O: 64.000 ORIFICE PRESSURE DROP, psi: 2.308 LIQUID DENSITY, 1b/ft^3: 42.260
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ORIFICE PRESSURE DROP, psi:2.308LIQUID DENSITY, lb/ft^3:42.260
LIQUID DENSITY, 1b/ft ³ : 42.260
LIQUID VISCOSITY, CP: .6000
LIQUID VOLUMETRIC RATE, ft^3/min.: 12.840
LIQUID MASS RATE, 1b/sec.: 9.044
ORIFICE SIZE, inch: 1.501
REYNOLDS NUMBER: 176582.

Table 5-8

Orifice Sizing for Gas Flow: Program Inputs and Outputs

```
DATA51.DAT
GAS
INCH-H2O
4.026 14.7
1.4 0.075
0.0219
20.0 0.25
                         ORIFICE SIZING FOR GAS FLOWS
TYPE OF FLOW:
                                                                      GAS
    TYPE OF PRESSURE DROP:
                                                                      INCH-H2O
    INSIDE PIPE DIAMETER, inch:
                                                                        4.026
    UPSTREAM PRESSURE, psia.:
ORIFICE PRESSURE DROP, INCH-H2O:
ORIFICE PRESSURE DROP, psi.:
SPECIFIC HEAT RATIO (k=Cp/Cv):
                                                                       14.700
                                                                       20.000
                                                                          .721
                                                                       1.4000
    UPSTREAM FLUID DENSITY, 1b/ft^3:
                                                                          .075
    GAS VISCOSITY, CP:
                                                                         .0219
    GAS FLOW RATE, ft^3/min.:
GAS MASS RATE, lb/sec.:
                                                                       200.000
                                                                         .250
    EXPANSION FACTOR:
                                                                         .0409
    ORIFICE DIAMETER, inch:
                                                                         1.811
    FULL SCALE GAS FLOWRATE, 1b/sec.:
                                                                          .250
    REYNOLDS NUMBER:
                                                                            64410.
```

the orifice diameter at the liquid flow rate of 12.84 ft³/min and pipe diameter of 1.939 in. is 1.501 in. For the gas flow rate of 0.25 lb/s and pipe internal diameter of 4.026 in., the orifice size is 1.811 in.

Problem 5.2

(1) Determine the control valve capacity coefficient, C_v , for the following liquid flow conditions:

Flow = 400 gpm $P_1 = 150 \text{ psia}$ $P_2 = 125 \text{ psia}$ $\text{SpGr}_1 = 1.0$

(2) Calculate the control valve coefficient for gas flow under the following conditions:

Flow = 1200 ft³/h P₁ = 550 psia P₂ = 150 psia SpGr_g = 0.6 T = 100°F

(3) Calculate the steam flow through a valve with a capacity coefficient, $C_v = 75$, under the following conditions:

°F superheat = 0 $P_1 = 70$ psia $P_2 = 60$ psia

Solutions

The computer program PROG52 calculates the control valve capacity coefficient (C_v) or the maximum flow rate that will pass through the valve. Table 5-9 shows the input data and results of the control valve calculations for liquid, gas, and steam conditions. For the liquid service, at a flow rate of 400 gpm and the control valve pressure drop of 25 psi, the control valve capacity coefficient (C_v) is 80. The gas service with a flow rate of 1200 ft³/h, and control valve pressure drop of 400 psi gives the control valve capacity coefficient (C_v) of 0.059. For the steam condition of capacity coefficient (C_v) = 75, and the control valve pressure drop of 10 psi, the maximum steam flow rate through the valve is 5511.4 lb/hr.

Problem 5.3

(1) In a biological plant, ammonia is used to control the pH during fermentation to obtain maximum yield of the product. For this purpose, liquid ammonia is vaporized by passing steam through a coil in a vaporizer. A relief valve from the vaporizer is set at a pressure of 290 psig. If the relieving rate of ammonia vapor is 620 lb/hr, determine the size of the relief valve required to relieve ammonia vapor during an emergency.

Design data are:

Ratio of specific heat capacities (C_p/C_v)	= 1.33
Molecular weight of ammonia	= 17.03
Critical Pressure, P _c	= 111.3 atm
Critical Temperature, T _c	= 405.6 K
Constants in Antoine's equation: A	= 16.9481
В	= 2132.5
С	= -32.98

Table 5-9 Control Valve Sizing for Liquid, Gas and Steam Conditions: Program Inputs and Outputs

DATA52.DAT	
LIQUID	
400.0 0.0	
150.0 125.0 1.0	
GAS	
1200.0 0.0	
550.0 150.0 0.6 100.0	
STEAM	
75.0	
70.0 60.0 0.0	
CONTROL VALVE SIZING FOR LIO	JID SERVICE

FLUID SERVICE:	LIQUID
LIQUID FLOWRATE, gpm:	400.00
LIQUID SPECIFIC GRAVITY:	1.000
INLET PRESSURE, psia:	150.000
OUTLET PRESSURE, psia:	125.000
CONTROL VALVE PRESSURE DROP, psi:	25.000
	25.000 80.000
CONTROL VALVE PRESSURE DROP, psi: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR GAS	80.000 SERVICE
CONTROL VALVE PRESSURE DROP, psi: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR GAS	80.000 SERVICE GAS
CONTROL VALVE PRESSURE DROP, psi: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR GAS FLUID SERVICE: GAS FLOW RATE, ft^3/hr.:	80.000 SERVICE GAS 1200.000
CONTROL VALVE PRESSURE DROP, psi: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR GAS FLUID SERVICE: GAS FLOW RATE, ft^3/hr.: SPECIFIC GRAVITY:	80.000 SERVICE GAS 1200.000 .600
CONTROL VALVE PRESSURE DROP, psi: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR GAS CONTROL VALVE SIZING FOR GAS FLUID SERVICE: GAS FLOW RATE, ft^3/hr.: SPECIFIC GRAVITY: INLET PRESSURE, psia:	80.000 SERVICE GAS 1200.000 600 550.000
CONTROL VALVE PRESSURE DROP, psi: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR GAS CONTROL VALVE SIZING FOR GAS FLUID SERVICE: GAS FLOW RATE, ft^3/hr.: SPECIFIC GRAVITY: INLET PRESSURE, psia: OULET PRESSURE, psia:	80.000 SERVICE GAS 1200.000 .600 550.000 150.000
CONTROL VALVE PRESSURE DROP, psi: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR GAS FLUID SERVICE: GAS FLOW RATE, ft^3/hr.: SPECIFIC GRAVITY: INLET PRESSURE, psia: OULET PRESSURE, psia: CONTROL VALVE PRESSURE DROP, psi:	80.000 SERVICE GAS 1200.000 .600 550.000 150.000 400.000
CONTROL VALVE PRESSURE DROP, psi: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR GAS CONTROL VALVE SIZING FOR GAS FLUID SERVICE: GAS FLOW RATE, ft^3/hr.: SPECIFIC GRAVITY: INLET PRESSURE, psia: OULET PRESSURE, psia:	80.000 SERVICE GAS 1200.000 .600 550.000 150.000
CONTROL VALVE PRESSURE DROP, psi: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR GAS CONTROL VALVE SIZING FOR GAS FLOW RATE, ft^3/hr.: SPECIFIC GRAVITY: INLET PRESSURE, psia: OULET PRESSURE, psia: CONTROL VALVE PRESSURE DROP, psi: INLET TEMPERATURE, oF: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE CAPACITY COEFFICIENT:	80.000 SERVICE GAS 1200.000 .600 550.000 150.000 400.000 100.000 .059
CONTROL VALVE PRESSURE DROP, psi: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR GAS FLUID SERVICE: GAS FLOW RATE, ft^3/hr.: SPECIFIC GRAVITY: INLET PRESSURE, psia: OULET PRESSURE, psia: CONTROL VALVE PRESSURE DROP, psi: INLET TEMPERATURE, oF: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE CAPACITY COEFFICIENT:	80.000 SERVICE GAS 1200.000 .600 550.000 150.000 400.000 100.000 .059 MM SERVICE
CONTROL VALVE PRESSURE DROP, psi: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR GAS FLUID SERVICE: GAS FLOW RATE, ft^3/hr.: SPECIFIC GRAVITY: INLET PRESSURE, psia: OULET PRESSURE, psia: CONTROL VALVE PRESSURE DROP, psi: INLET TEMPERATURE, oF: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR STEA FLUID SERVICE:	80.000 SERVICE GAS 1200.000 550.000 150.000 400.000 100.000 .059
CONTROL VALVE PRESSURE DROP, psi: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR GAS FLUE RATE, ft^3/hr.: SPECIFIC GRAVITY: INLET PRESSURE, psia: OULET PRESSURE, psia: CONTROL VALVE PRESSURE DROP, psi: INLET TEMPERATURE, oF: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR STEL FLUID SERVICE: CALCULATED STEAM FLOW RATE, 1b/hr.:	80.000 SERVICE GAS 1200.000 .600 550.000 150.000 400.000 100.000 .059 MM SERVICE STEAM 5511.352
CONTROL VALVE PRESSURE DROP, psi: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR GAS FLUID SERVICE: GAS FLOW RATE, ft^3/hr.: SPECIFIC GRAVITY: INLET PRESSURE, psia: OULET PRESSURE, psia: CONTROL VALVE PRESSURE DROP, psi: INLET TEMPERATURE, OF: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR STEA FLUID SERVICE: CALCULATED STEAM FLOW RATE, lb/hr.: DEGREE OF SUPERHEAT, oF:	80.000 SERVICE GAS 1200.000 .600 550.000 150.000 400.000 100.000 .059 MM SERVICE STEAM 5511.352 .000
CONTROL VALVE PRESSURE DROP, psi: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR GAS FLUID SERVICE: GAS FLOW RATE, ft^3/hr.: SPECIFIC GRAVITY: INLET PRESSURE, psia: OULET PRESSURE, psia: CONTROL VALVE PRESSURE DROP, psi: INLET TEMPERATURE, oF: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE CAPACITY COEFFICIENT: FLUID SERVICE: CALCULATED STEAM FLOW RATE, lb/hr.: DEGREE OF SUPERHEAT, oF: INLET PRESSURE, psia:	80.000 SERVICE GAS 1200.000 550.000 150.000 400.000 100.000 .059 MM SERVICE STEAM 5511.352 .000 70.000
CONTROL VALVE PRESSURE DROP, psi: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR GAS FLUID SERVICE: GAS FLOW RATE, ft^3/hr.: SPECIFIC GRAVITY: INLET PRESSURE, psia: OULET PRESSURE, psia: CONTROL VALVE PRESSURE DROP, psi: INLET TEMPERATURE, OF: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE CAPACITY COEFFICIENT: CONTROL VALVE SIZING FOR STEA FLUID SERVICE: CALCULATED STEAM FLOW RATE, lb/hr.: DEGREE OF SUPERHEAT, oF:	80.000 SERVICE GAS 1200.000 .600 550.000 150.000 400.000 100.000 .059 MM SERVICE STEAM 5511.352 .000

(2) A bellow style pressure relief valve is required to protect a vessel containing an organic liquid. The required relieving capacity is 310 gpm. Inlet temperature is 170°F. Set pressure is 100 psig. Allowable overpressure is 25%. Built-up back pressure is 25 psig. Specific gravity is 1.45 and viscosity is 3,200 cP. Determine the orifice size of the valve. The correction factors are:

 $K_v = 1.0$ $K_w = 0.925$ $K_d = 0.65$

(3) Calculate the required area for the relief valve for a horizontal, noninsulated storage tank exposed to fire and containing liquid vinyl chloride monomer. The tank dimensions are shown in Figure 5-14, with the design pressure of 100 psig. Discharge from the relief valve will be vented to a gas holder operating at 0.5 psig. A 20% accumulation is assumed over the design pressure. Average inventory of the tank contents will equal 75% of the vessel's inside diameter.

Solutions

The computer program PROG53 sizes relief valves for vapor, steam, liquid, air, and fire conditions. Tables 5-10 and 5-11 show the input data and results of the vapor and liquid services. The results of the relief valve sizing of an ammonia vaporizer at the relieving rate of 620 lb/hr and a set pressure of 290 psig, indicate that the calculated orifice area is 0.030 in². The nearest standard orifice size is D with an orifice area of 0.110 in². The relief valve calculations for the organic liquid at the relieving rate of 310 gpm and a set pressure of 100 psig give the calculated orifice area of 2.233 in². The nearest standard orifice size is L, with an orifice area of 2.853 in².

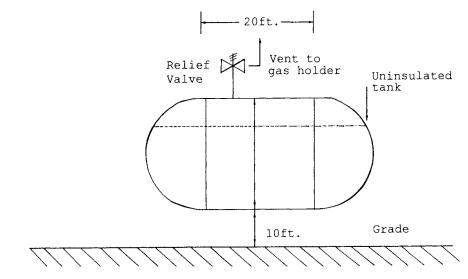


Figure 5-14. A horizontal, non-insulated storage tank.

Table 5-10 Relief Valve Sizing for Vapor Service: Program Inputs and Outputs

DATA53.DAT	
VAPOR	
620.0 290.0	
10.0 133.4	
0.84 17.03	
1.31	
VAPOR RELIEF VALVE SIZING	
***************************************	*****
BASIS OF SELECTION:	VAPOR
REQUIRED VAPOR RATE, lb/hr.:	620.
VALVE SET PRESSURE, psig:	290.000
PERCENTAGE IN OVERPRESSURE, (%):	10.000
UPSTREAM FLUID TEMPERATURE, DEG. F:	133.400
COMPRESSIBILITY FACTOR:	.8400
MOLECULAR WEIGHT OF VAPOR:	17.030
RATIO OF SPECIFIC HEATS OF FLUID:	1.310
GAS CONSTANT DETERMINED BY RATIO OF Cp/Cv:	347.913
CALCULATED VALVE ORIFICE AREA, in^2:	.030
NEAREST STANDARD ORIFICE AREA, in^2:	.110
NEAREST STANDARD ORIFICE SIZE:	D
PREFERRED VALVE BODY SIZE: (IN-OUT) inches:	1-2
MAXIMUM VAPOR FLOWRATE WITH STD. ORIFICE, lb/hr.:	2302.
REACTION FORCE, 1b:	28.

 Table 5-11

 Relief Valve Sizing for Liquid Service: Program Inputs and Outputs

I

DATA53.DAT	
LIQUID	
310.0 100.0	
25.0 1.45	
3200.0 1.0	
0.925 0.65	
RELIEF VALVE SIZING FOR LIQUID	

BASIS OF SELECTION:	LIQUID
LIQUID FLOWRATE, US gpm:	310.
VALVE SET PRESSURE, psig:	100.000
BACK PRESSURE ON VALVE, psig: FluID SPECIFIC GRAVITY:	25.000
FLUID VISCOSITY, CP:	1.450
	3200.0000
VISCOSITY CORRECTION FACTOR, KV: BACKPRESSURE CORRECTION FACTOR, KW:	1.0000
EFFECTIVE COEFFICIENT OF DISCHARGE, Kd:	.9250
CALCULATED VALVE ORIFICE AREA, in^2:	.650
NEAREST STANDARD ORIFICE AREA, in^2:	1.772
NEAREST STANDARD ORIFICE AREA, 11-2: NEAREST STANDARD ORIFICE SIZE:	1.838
PREFERRED VALVE BODY SIZE, (IN-OUT) inches:	ĸ
MAXIMUM LIQUID FLOWRATE WITH STD. ORIFICE, qpm:	3-4
VISCOSITY CORRECTION FACTOR:	322.
REYNOLDS NUMBER:	.794
CALCULATED VALVE ORIFICE AREA, in^2:	290.
NEAREST STANDARD ORIFICE AREA, in^2:	2.233
NEAREST STANDARD ORIFICE SIZE:	2.853 L
PREFERRED VALVE BODY SIZE, (IN-OUT) inches:	
MAXIMUM LIQUID FLOWRATE WITH STD. ORIFICE, gpm:	3-4 or 4-6 396.

Fire Relief Valve Sizing

The wetted surface for the vessel equals the wetted area of the two heads plus that of the cylindrical section. Thus:

$$A_{w} = \frac{2\pi (yD_{1})^{2}}{4} + D_{2}L\pi$$

where y = the fraction of the vessel's internal diameter (ID) that is equivalent to average liquid inventory.

 D_1 = diameter of circular blank from which head is shaped (D_1 will depend on the type of head).

 $D_2 = y$ times the ID of the tank.

L = tangent-to-tangent length of the cylindrical section.

$$D_1 = 11 \text{ ft}$$

Therefore, the wetted surface area is:

$$A_{w} = \frac{2\pi [(0.75)(11)]^{2}}{4} + \pi (0.75)(10)(20)$$
$$A_{w} = 578.15 \text{ ft}^{2}$$

The relieving temperature at the set pressure plus 20% accumulation plus 14.7 psia = 135° F. At this temperature, the latent heat of vaporization is 116 Btu/lb. Molecular weight of vinyl chloride, $M_w = 62.5$. Ratio of specific heat capacities, k = 1.17. The critical properties of vinyl chloride are:

 $P_c = 809 \text{ psia}$ $T_c = 313.7^{\circ}\text{F}$ $P_r = P/P_c = 134.7/809 = 0.165$ $T_r = T/T_c = (135 + 460)/(313.7 + 460) = 0.77$

From a compressibility factor chart, Z = 0.86. The rate of heat input due to fire and flow rate of the vapor release can be determined by Equations 5-53 and 5-45. Table 5-12 gives the input data and results of vinyl chloride monomer. The results show that the calculated relief valve orifice area is 2.172 in². The nearest standard orifice size is L, with an orifice area of 2.853 in².

 Table 5-12

 Relief Valve Sizing for Fire Service: Program Inputs and Outputs

DATA53.DAT	
 FIRE	
1.0 578.15	
116.0 100.0	
20.0 135.0	
0.86 62.5	
1.17	
FIRE RELIEF VALVE SIZING	
***************************************	******
BASIS OF SELECTION:	FIRE
ENVIRONMENTAL FACTOR:	1.000
WETTED SURFACE AREA OF VESSEL, ft^2:	578.150
LATENT HEAT OF VAPORIZATION, Btu/lb:	116.000
VALVE SET PRESSURE, psig:	100.000
PERCENTAGE OF OVERPRESSURE, (%):	20.000
FLUID TEMPERATURE, DEG. F:	135.000
FLUID COMPRESSIBILITY FACTOR:	.860
FLUID MOLECULAR WEIGHT:	62.500
RATIO OF SPECIFIC HEATS OF FLUID:	1,170
GAS CONSTANT DETERMINED BY RATIO OF Cp/Cv:	334.173
VAPOR RELIEF FLOW RATE, 1b/h.:	33315.
CALCULATED VALVE ORIFICE AREA, in^2:	2.172
NEAREST STANDARD ORIFICE AREA, in^2:	2.853
NEAREST STANDARD ORIFICE SIZE:	L
PREFERRED VALVE BODY SIZE, (IN-OUT) inches:	3-4 or 4-6
MAXIMUM VAPOR FLOWRATE WITH STD. ORIFICE 1b/hr.:	43760.
REACTION FORCE, 1b:	271.

Problem 5.4

Determine the vent size for a vapor pressure system using the following data and physical properties.

Reactor Parameters

Volume, V = 10 m³ Mass, M = 8,000 kg Vent opening pressure = 15 bara = 217.5 psia Temperature at set pressure $T_s = 170^{\circ}C$ = 443.15K Overpressure allowed above operating pressure = 10% = 1bar (10⁵ Pa)

Material Properties

Specific heat, $C_p = 3000 \text{ J/kg.K}$ Slope of vapor pressure and temperature curve dP/dT = 20,000 Pa/K

Rate of Reaction

 $\Delta T = dP/20,000 = 10^{5}/20,000 = 5K$ Rate at set pressure, $(dT/dt)_{s} = 6.0$ K/min Rate at maximum pressure = 6.6 K/min Average rate = 6.3 K/min = 0.105 K/s

Solution

Using Fauske's nomograph of Figure 5-12 at a self heat rate of 6.3 K/ min. and a set pressure of 217.5 psia, the corresponding vent area per 1,000 kg of reactants = 0.0008m^2 .

The vent area of 8,000 kg reactants = 0.0064m² The vent size

d =
$$\left(\frac{4 \text{Area}}{\pi}\right)^{0.5}$$
 = 90.27 mm (3.6 in.)

If Figure 5-12 is applicable for F=0.5, then for F=1.0, the area is

A =
$$(0.0064 \,\mathrm{m}^2) \left(\frac{0.5}{1.0}\right) = 0.0032 \,\mathrm{m}^2$$

The area assumes a 20% absolute overpressure.

The result can be adjusted for other overpressures by multiplying the area by a ratio of 20/(new absolute percent overpressure).

Using the Leung's method:

Assuming L/D = 0, F = 1.0 Two-phase mass flux from Equation 5-58

G = (0.9)(1.0)(20,000)
$$\left(\frac{1.0 \times 443.15}{3,000}\right)^{0.5}$$

$$= 6918.1 \frac{\text{kg}}{\text{m.s}^2}$$

Rate of heat generation: q

q = C_p
$$\frac{dT}{dt} \left(\frac{J.K}{kg.K.s} \right) = (3,000)(0.105) \frac{W}{kg} = 315 \frac{W}{kg}$$

From Equation 5-63, the vent area, A, is:

A =
$$\frac{8,000 \times 315}{6918.1 \left\{ \left(\frac{10}{8,000} \times 443.15 \times 20,000 \right)^{0.5} + (3,000 \times 5.0)^{0.5} \right\}^2}$$

A = 7.024 × 10⁻³ m²
d = 94.56 mm (3.72 in.)

The vent size is about 4.0 in. Using the Fauske's method:

Assuming $\alpha = 1.0, \alpha_0 = 0.0, F = 1.0$

 $\Delta P = 1 \text{ bar} = 10^5 \text{N/m}^2$

From Equation 5-68, the vent area, A, is

A =
$$\frac{1}{2} \bullet \frac{(8,000)(0.105)(1-0)}{(1.0)\left(\frac{443.15}{3,000}\right)^{0.5}(10^5)(1-0)}$$

$$A = 10.927 \times 10^{-3} \,\mathrm{m}^2$$

d = 117.95 mm(4.64 in.)

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PROGRAM PROG51

-			
с	ORIFICE SIZING FOR FLUID FLOWS.		
с	THIS PROGRAM CALCULATES AN ORIFICE DIAMETER FOR BOTH LIQUID AND GAS		
с	FLOWS USING FULL-SCALE FLOWRATES IN EITHER 1b/s. OR ft^3/min. AND THE		
с	FULL-SCALE PRESSURE DROPS IN EITHER PSI. OR INCH-H2O.		
с	THE PROGRAM APPLIES TO CORNER, FLANGE, VENA-CONTRACTA, AND		
с	1/2 DIAMETER TAPS.		
č			**********************
č			
č	ACFM	=	FULL-SCALE FLOWRATE AT UPSTREAM CONDITIONS, ft^3/min.
с	CD	=	
с	D	=	
с	OD	=	
с	FA	=	AREA THERMAL-EXPANSION FACTOR. ACCOUNTS FOR THE
с			EXPANSION OF THE ORIFICE WITH TEMPERATURE CHANGES.
с	Hw	=	FULL-SCALE ORIFICE PRESSURE-DIFFERENTIAL. INCH-H20 AT
č			680F.
č	к	-	
č	G	-	
с	UP	=	orbridder inddetond, perti
с	DELP	=	FULL-SCALE ORIFICE PRESSURE-DIFFERENTIAL, psi.
с	Y	=	EXPANSION FACTOR.
с	BETA	=	RATIO OF ORIFICE DIAMETER TO PIPE INTERNAL DIAMETER.
с	DEN	-	UPSTREAM FLUID DENSITY, 1b/ft^3.
c	CHOICE		
č	0.10101		OR INCH-H2O.
c	FLOW	-	CHECKS WHETHER THE FLOW CONDITION IS LIQUID OR GAS.
c			**************************************
C	******	****	***************************************
	CHARACT	ER*1	0 CHOICE, FLOW
			O CHOICE, FLOW OD, N, L, FMASS
	REAL K,	ID,	
	REAL K,	ID,	OD, N, L, FMASS
	REAL K, COMMON/	ID, DATA	OD, N, L, FMASS 1/FLOW, CHOICE
	REAL K, COMMON/ OPEN (U	ID, DATA	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18)
	REAL K, COMMON/ OPEN (U	ID, DATA	OD, N, L, FMASS 1/FLOW, CHOICE
G	REAL K, COMMON/ OPEN (U OPEN (U	ID, DATA NIT= NIT=	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1, FILE ='PRN')
c	REAL K, COMMON/ OPEN (U OPEN (U CHECK W	ID, DATA NIT= NIT= NIT=	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18)
C C	REAL K, COMMON/ OPEN (U OPEN (U	ID, DATA NIT= NIT= NIT=	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1, FILE ='PRN')
с	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2	ID, DATA NIT= NIT= HETH O.	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1, FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR
	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2	ID, DATA NIT= NIT= HETH O.	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1, FILE ='PRN')
с	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2 CHOOSE	ID, DATA NIT= NIT= HETH O. TYPE	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1, FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR TOF FLOW FOR SIZING (LIQUID/GAS)
с	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2 CHOOSE	ID, DATA NIT= NIT= HETH O. TYPE	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1, FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR
с	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2 CHOOSE	ID, DATA NIT= NIT= HETH O. TYPE	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1, FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR TOF FLOW FOR SIZING (LIQUID/GAS)
c c	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2 CHOOSE READ (3	ID, DATA NIT= NIT= HETH O. TYPE	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1, FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR TOF FLOW FOR SIZING (LIQUID/GAS)
c c	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2 CHOOSE READ (3 FORMAT	ID, DATA NIT= NIT= HETH O. TYPE (A)	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1, FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR OF FLOW FOR SIZING (LIQUID/GAS) ERR=19) FLOW
c c	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2 CHOOSE READ (3 FORMAT	ID, DATA NIT= NIT= HETH O. TYPE (A)	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1, FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR TOF FLOW FOR SIZING (LIQUID/GAS)
c c	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2 CHOOSE READ (3 FORMAT IF (FLO	ID, DATA NIT= NIT= HETH O. TYPE (A) (A)	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1, FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR OF FLOW FOR SIZING (LIQUID/GAS) ERR=19) FLOW
c c	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2 CHOOSE READ (3 FORMAT	ID, DATA NIT= NIT= HETH O. TYPE (A) (A)	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1, FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR OF FLOW FOR SIZING (LIQUID/GAS) ERR=19) FLOW
c c	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2 CHOOSE READ (3 FORMAT IF (FLO GO TO 2	ID, DATA NIT= NIT= HETH O. TYPE (A) W .E	OD, N, L, FMASS 1/FLOW, CHOICE 3. FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1. FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR OF FLOW FOR SIZING (LIQUID/GAS) ERR=19) FLOW Q. 'LIQUID' .OR. FLOW .EQ. 'liquid') THEN
c c	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2 CHOOSE READ (3 FORMAT IF (FLO GO TO 2 ELSEIF	ID, DATA NIT= NIT= HETH O. TYPE (A) W .E	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1, FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR OF FLOW FOR SIZING (LIQUID/GAS) ERR=19) FLOW
c c	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2 CHOOSE READ (3 FORMAT IF (FLO GO TO 2	ID, DATA NIT= NIT= HETH O. TYPE (A) W .E	OD, N, L, FMASS 1/FLOW, CHOICE 3. FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1. FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR OF FLOW FOR SIZING (LIQUID/GAS) ERR=19) FLOW Q. 'LIQUID' .OR. FLOW .EQ. 'liquid') THEN
С С 5	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2 CHOOSE READ (3 FORMAT IF (FLO GO TO 2 ELSEIF ENDIF	ID, DATA NIT= NIT= TYPE (A) (A) (FLC	OD, N, L, FMASS 1/FLOW, CHOICE 3. FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1. FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR OF FLOW FOR SIZING (LIQUID/GAS) ERR=19) FLOW Q. 'LIQUID' .OR. FLOW .EQ. 'liquid') THEN W .EQ. 'GAS' .OR. FLOW .EQ. 'gas') THEN
c c	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2 CHOOSE READ (3 FORMAT IF (FLO GO TO 2 ELSEIF ENDIF	ID, DATA NIT= NIT= TYPE (A) (A) (FLC	OD, N, L, FMASS 1/FLOW, CHOICE 3. FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1. FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR OF FLOW FOR SIZING (LIQUID/GAS) ERR=19) FLOW Q. 'LIQUID' .OR. FLOW .EQ. 'liquid') THEN
С С 5	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2 CHOOSE READ (3 FORMAT IF (FLO GO TO 2 ELSEIF ENDIF CHOOSE	ID, DATA NIT= NIT= HETH O. TYPE 0 (FLC TYPE	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1, FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR C OF FLOW FOR SIZING (LIQUID/GAS) ERR=19) FLOW Q. 'LIQUID' .OR. FLOW .EQ. 'liquid') THEN W .EQ. 'GAS' .OR. FLOW .EQ. 'gas') THEN OF PRESSURE DROP.
С С 5	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2 CHOOSE READ (3 FORMAT IF (FLO GO TO 2 ELSEIF ENDIF CHOOSE	ID, DATA NIT= NIT= HETH O. TYPE 0 (FLC TYPE	OD, N, L, FMASS 1/FLOW, CHOICE 3. FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1. FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR OF FLOW FOR SIZING (LIQUID/GAS) ERR=19) FLOW Q. 'LIQUID' .OR. FLOW .EQ. 'liquid') THEN W .EQ. 'GAS' .OR. FLOW .EQ. 'gas') THEN
С С 5	REAL K, COMMON/ OPEN (U OPEN (U CHECK W INCH-H2 CHOOSE READ (3 FORMAT IF (FLO GO TO 2 ELSEIF ENDIF CHOOSE	ID, 'DATA NIT= NIT= ''NIT= ''NIT=	OD, N, L, FMASS 1/FLOW, CHOICE 3, FILE ='DATA51.DAT', STATUS ='OLD', ERR=18) 1, FILE ='PRN') ER THE DATA FOR THE ORIFICE PRESSURE DROP IS PSI. OR C OF FLOW FOR SIZING (LIQUID/GAS) ERR=19) FLOW Q. 'LIQUID' .OR. FLOW .EQ. 'liquid') THEN W .EQ. 'GAS' .OR. FLOW .EQ. 'gas') THEN OF PRESSURE DROP.

с с с с с с	D = PIPE INTERNAL DIAMETER, inch. UP = THE UPSTREAM PRESSURE, psia. K = THE RATIO OF SPECIFIC HEAT CAPACITIES, Cp/Cv. DEN = THE GAS DENSITY, lb/ft^3. VISG = THE GAS VISCOSITY, cP.
	READ (3, *, ERR=19) D, UP READ (3, *, ERR=19) K, DEN READ (3, *, ERR=19) VISG
	IF (CHOICE .EQ. 'INCH-H2O' .OR. CHOICE .EQ. 'inch-H2O') THEN
c c	READ THE ORIFICE PRESSURE DROP IN INCH-H2O AND THE FLUID FLOW RATE IN $1b/s$.
	READ (3, *, ERR=19) DELP2, G2
c c	CALCULATE THE ORIFICE PRESSURE DROP IN PSI. CALCULATE THE GAS FLOW RATE IN ft^3/min.
	DELP1=0.036063*DELP2 G1 = (G2*60.0)/DEN
	GO TO 25
	ELSEIF (CHOICE .EQ. 'PSI' .OR. CHOICE .EQ. 'psi') THEN
c c	READ THE ORIFICE PRESSURE DROP IN PSI. AND THE GAS FLOW RATE IN ft^3/min.
	READ (3, *, ERR=19) DELP1, G1
c c	GAS FLOWRATE, (G2) IN lb/sec. ORIFICE PRESSURE DROP (DELP2) IN INCH-H20
	G2 = (G1*DEN)/60.0 DELP2 = 27.72925*DELP1 ENDIF
	GO TO 25
20 30	READ (3, 30, ERR=19) CHOICE FORMAT (A)
	IF (CHOICE .EQ. 'INCH-H2O' .OR. CHOICE .EQ. 'inch-H2O') THEN
с с с с с с	ID = PIPE INTERNAL DIAMETER, inch. DELP = ORIFICE PRESSURE DROP, INCH-H2O DEN = LIQUID DENSITY, lb/ft^3. Q = LIQUID FLOWRATE, ft^3/min. VISL = LIQUID VISCOSITY, cP.

	READ (3, *, ERR=19) ID, DELP READ (3, *, ERR=19) DEN, Q, VISL
c c	CALCULATE THE ORIFICE PRESSURE DROP IN PSI. CALCULATE THE LIQUID FLOW RATE IN lb/sec.
	DELP1 = 0.036063*DELP FMASS = (Q*DEN)/60.0
	CALL LIQUIDF (ID, DELP, DELP1, DEN, Q, FMASS, VISL)
	GO TO 999
	ELSEIF (CHOICE .EQ. 'PSI' .OR. CHOICE .EQ. 'psi') THEN
00000	<pre>ID = PIPE INTERNAL DIAMETER, inch. DELP1 = ORIFICE DIFFERENTIAL PRESSURE, PSI. DEN = LIQUID DENSITY, lb/ft.^3. FMASS = LIQUID FLOW RATE, lb/sec. VISL = LIQUID VISCOSITY, cP.</pre>
	READ (3, *, ERR=19) ID, DELP1 READ (3, *, ERR=19) DEN, FMASS, VISL
c c	CALCULATE THE ORIFICE PRESSURE DROP IN INCH-H20 CALCULATE THE LIQUID FLOW RATE IN ft^3/min.
	DELP = 27.72925*DELP1 Q = (FMASS*60.0)/DEN
	CALL LIQUIDF (ID, DELP, DELP1, DEN, Q, FMASS, VISL)
	ENDIF
	GO TO 999
18 100	WRITE (*, 100) Format (3x,'data file does not exist')
19 110	GO TO 999 WRITE (*, 110) FORMAT (3X,'ERROR MESSAGE IN THE DATA VALUE')
	GO TO 999
25	N = 1.0-(0.41*DELP1)/(UP*K) T = (0.35*DELP1)/(UP*K)
	SRT = (DEN *DELP1)**0.5 L = G2/(0.3255*(D**2)*SRT) P = (2.0*N)/T
	$Q = (N^{*}2+L^{*}2)/(T^{*}T)$

```
R = -(L + 2)/(T + T)
            A = (3.0*Q+(P*P))/3.0
            B = ((2.0*(P**3))-(9.0*P*Q)+(27.0*R))/27.0
            SUM1 = (((B**2)/4.0)+((A**3)/27.0))**0.5
            VAL1 = (-B/2.0+SUM1)**(1.0/3.0)
             Y = VAL1 + ((-B/2-SUM1)**(1.0/3.0))
            Y = ABS(Y-(2.0*N)/(3.0*T))
            OD = D*(Y**0.25)
            USING THE ORIFICE DIAMETER, CHECK WHETHER THE CALCULATED GAS
RATE IS THE SAME AS THE INITIAL GAS RATE.
c
c
с
            CALCULATE THE EXPANSION FACTOR, Y1
            BETA = OD/D
            Y1 = 1.0-(0.41+0.35*(BETA**4))*(DELP1/(UP*K))
            FA = 1.0
            CD = 0.62
            VAL1 = (1.0 - (BETA * * 4)) * * 0.5
            G = 0.52502 * CD * Y1 * (OD * * 2) * FA * SRT
            GR = G/VAL1
с
            CALCULATE THE REYNOLDS NUMBER
            REG = (6.31*3600.0*G2)/(D*VISG)
            WRITE (1, 120)
 120
            FORMAT (25X, 'ORIFICE SIZING FOR GAS FLOWS', /1H ,76(1H*))
            WRITE (1, 130)FLOW
 130
            FORMAT (5X, 'TYPE OF FLOW: ', T65, A)
            WRITE (1, 140) CHOICE
 140
            FORMAT (5X, 'TYPE OF PRESSURE DROP:', T65, A)
            WRITE (1, 150) D, UP, DELP2, DELP1, K, DEN, VISG, G1, G2
            WRITE (1, 150) D, UP, DEEP2, DELP1, K, DEN, VISG, GI, G2
FORMAT (5X,'INSIDE PIPE DIAMETER, inch:', T65, F8.3,/,
5X,'UPSTREAM PRESSURE, psia.:', T65, F8.3,/,
5X,'ORIFICE PRESSURE DROP, INCH-H20:', T65, F8.3,/,
5X,'SPECIFIC HEAT RATIO (k=Cp/Cv):', T65, F8.4,/,
5X,'UPSTREAM FLUID DENSITY, lb/ft^3:', T65, F8.3,/,
5Y,'G5 VISCOSITY op:' T65 F8.4,/,
5Y,'G5 VISCOSITY op:' T65 F8.4,/,
 150
        *
        *
                         5X,'GAS VISCOSITY, CP:', T65, F8.4,/,
5X,'GAS FLOW RATE, ft^3/min.:', T65, F8.3,/,
5X,'GAS MASS RATE, lb/sec.:', T65, F8.3)
        *
        ٠
            WRITE (1, 160)Y, OD, GR, REG
            WRITE (1, 100)1, 00, GR, REG
FORMAT (5X,'EXPANSION FACTOR:', T65, F8.4,/,
5X,'ORIFICE DIAMETER, inch:', T65, F8.3,/,
5X,'FULL SCALE GAS FLOWRATE, lb/sec.:', T65, F8.3,/,
 160
        ٠
```

```
5X, 'REYNOLDS NUMBER:', T65, F12.0)
    *
        WRITE (1, 170)
170
        FORMAT (1H ,76(1H-))
        FORM FEED THE PRINTING PAPER TO THE TOP OF NEXT PAGE.
C
        WRITE (1, *) CHAR(12)
        CLOSE (UNIT=3, STATUS='KEEP')
        CLOSE (1)
999
        STOP
        END
с
с
с
с
        ****
        THE FORMULA FOR CALCULATING THE ORIFICE PLATE FOR LIQUID
        FLOW IS:
С
С
С
С
        M = 0.52502((Cd.Y.d^{2}.Fa)(SQRT(DEN.DELP)/(SQRT(1-(DO/D)^{4})))
        WHERE FOR LIQUID FLOW, Cd=0.62, Fa=1.0 AND Y=1.0
            = ORIFICE DIAMETER
        DO
с
с
с
с
        D
             = PIPE INTERNAL DIAMETER
        DEN = DENSITY OF LIQUID, 1b/ft^3.
        DELP = PRESSURE DROP ACROSS THE ORIFICE, inch-H20 OR psi.
            = MASS FLOW RATE, lb/s.
        м
с
        ********
                             ...
                                ************************************
        SUBROUTINE LIQUIDF (ID, DELP, DELP1, DEN, Q, FMASS, VISL)
        CHARACTER*10 FLOW, CHOICE
        COMMON/DATA1/FLOW, CHOICE
        REAL ID, OD
        CALCULATE THE ORIFICE DIAMETER, inch.
С
        PRESSURE DROP ACROSS THE ORIFICE (DELP1), PSI.
С
с
        LIQUID FLOW RATE (FMASS), lb/sec.
        Z = (0.52502*0.62*(DEN*DELP1)**0.5)/FMASS
        RATIO = (ID**4)/(1+((ID**4)*(Z**2)))
        OD = RATIO**0.25
        CALCULATE THE REYNOLDS NUMBER, REL
С
        REL = (6.31*3600.0*FMASS)/(ID*VISL)
с
        PRINT THE RESULTS OF ORIFICE SIZING FOR LIQUID FLOW.
        WRITE (1, 180)
        FORMAT (20X, 'ORIFICE SIZING FOR LIQUID FLOW', /1H ,76(1H*))
180
        WRITE (1, 190) FLOW
190
        FORMAT (5X, 'TYPE OF FLOW:', T65, A)
        WRITE (1, 200)CHOICE
 200
        FORMAT (5X, 'TYPE OF PRESSURE DROP: ', T65, A)
```

210 * * * * *	5X, 'ORIFICE PRESSURE DROP, inch-H20:',T65,F8.3,/, 5X,'ORIFICE PRESSURE DROP, psi:', T65,F8.3,/, 5X,'LIQUID DENSITY, lb/ft^3:', T65,F8.3,/, 5X,'LIQUID VISCOSITY, cP:', T65,F8.4,/, 5X,'LIQUID VOLUMETRIC RATE, ft^3/min.:',T65,F8.3,/, 5X,'LIQUID MASS RATE, lb/sec.:', T65,F8.3,/,
220	WRITE (1, 220) Format (1H ,76(1H-))
с	FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR(12)
	RETURN

END

PROGRAM PROG52

0000000	THIS PROGRAM CALCULATES THE CONTROL VALVE CAPACITY COEFFICIENT (Cv) OR MAXIMUM FLUID RATE FOR LIQUID, VAPOR AND STEAM IN PROCESS OPERATIONS. IF THE CONTROL-VALVE COEFFICIENT(Cv) IS ALREADY KNOWN, THE PROGRAM WILL CALCULATE THE MAXIMUM FLUID FLOW RATE (LIQUID, VAPOR OR STEAM) THAT CAN PASS THROUGH THE VALVE FOR A GIVEN SET OF PROCESS CONDITIONS. TO COMPUTE THE FLUID FLOW FOR LIQUID, GAS AND STEAM CONDITIONS,
	<pre>VALUES OF THE VALVE CAPACITY COEFFICIENT MUST BE KNOWN (i.e. Cv>0) ************************************</pre>
с с с	***************************************
C	CHARACTER*10 CHOICE
	COMMON/DATA1/ V, P1, P2, DELPL, SPL, CVL COMMON/DATA2/ Q, P21, P22, DELPG, SPG, T, CVG COMMON/DATA3/ W, P31, P32, DELPS, F, CVS
	OPEN (UNIT=3, FILE='DATA52.DAT', STATUS='OLD', ERR=18) OPEN (UNIT=1, FILE='PRN')
5 6	READ (3, 6)CHOICE FORMAT (A)
	IF (CHOICE .EQ. 'LIQUID' .OR. CHOICE .EQ. 'liquid') THEN GO TO 11 ELSEIF (CHOICE .EQ. 'GAS' .OR. CHOICE .EQ. 'gas') THEN GO TO 22 ELSEIF (CHOICE .EQ. 'STEAM' .OR. CHOICE .EQ. 'steam') THEN GO TO 33 ENDIF
11	READ(3,*,ERR=10) V, CVL READ(3,*,ERR=10) P1, P2, SPL
	GO TO 44
22	READ(3,*,ERR=10) Q, CVG READ(3,*,ERR=10) P21, P22, SPG, T
	GO TO 55
33	READ(3,*,ERR=10) CVS

```
READ(3,*,ERR=10) P31, P32, F
            GO TO 66
 18
            WRITE(1,12)
 12
            FORMAT(3X, 'FILE DOES NOT EXIST')
 10
            WRITE(1,13)
            GO TO 999
  13
            FORMAT(3X,'ERROR MESSAGE IN THE DATA VALUE')
            GO TO 999
  44
            WRITE (1, 100)
            FORMAT (//,20X,'CONTROL VALVE SIZING FOR LIQUID SERVICE',/,
  100
                         1H ,76(1H*))
с
            CALCULATE THE CONTROL VALVE FOR LIQUID SERVICE
            CALL LIQUID
            WRITE (1, 110) CHOICE
            FORMAT (5X, 'FLUID SERVICE:', T60, A)
 110
            IF (CVL .GT. 0.0) THEN
            WRITE (1, 120) V, SPL, P1, P2, DELPL, CVL
 120
            FORMAT (5X,'LIQUID FLOWRATE, gpm:', T60, F9.2,/,
5X,'LIQUID SPECIFIC GRAVITY:', T60, F9.3,/,
                         5X, ILLET PRESSURE, psia:', T60, F9.3,/,
5X, COUTLET PRESSURE, psia:', T60, F9.3,/,
5X, CONTROL VALVE PRESSURE DROP, psi:', T60, F9.3,/,
5X, CONTROL VALVE CAPACITY COEFFICIENT:', T60, F9.3)
        *
        *
        ÷
            WRITE (1, 130)
 130
            FORMAT (76(1H-))
            ELSE
            WRITE (1, 140) V, SPL, P1, P2, DELPL, CVL
FORMAT (5X,'LIQUID FLOWRATE, gpm:', T60, F9.2,/,
5X,'LIQUID SPECIFIC GRAVITY:', T60, F9.3,/,
5X,'INLET PRESSURE, psia:', T60, F9.3,/,
5X,'CONTECT PRESSURE, psia:', T60, F9.3,/,
5X,'CONTROL VALVE PRESSURE DROP, psi:', T60, F9.3,/,
5X,'CONTROL VALVE CAPACITY COEFFICIENT:', T60, F9.3)
 140
        *
        *
        *
        .
            WRITE (1, 150)
FORMAT (76(1H-))
 150
            ENDIF
            GO TO 5
            WRITE (1, 160)
 55
 160
            FORMAT (//,20X,'CONTROL VALVE SIZING FOR GAS SERVICE',/,76(1H*))
С
            CALCULATE THE CONTROL VALVE FOR GAS SERVICE.
```

CALL GAS WRITE (1, 170) CHOICE 170 FORMAT (5X, 'FLUID SERVICE:', T60, A) IF (CVG .GT. 0.0) THEN WRITE (1, 180) Q, SPG, P21, P22, DELPG, T, CVG FORMAT (5X,'GAS FLOW RATE, ft^3/hr.:', T60, F9.3,/, 5X,'SPECIFIC GRAVITY:', T60, F9.3,/, 5X,'INLET PRESSURE, psia:', T60, F9.3,/, 5X,'OULET PRESSURE, psia:', T60, F9.3,/, 5X,'CONTROL VALVE PRESSURE DROP, psi:', T60, F9.3,/, 5X,'INLET TEMPERATURE, oF:', T60, F9.3,/, 5X,'CONTROL VALVE CAPACITY COEFFICIENT:', T60, F9.3) 180 * * * * ٠ ELSE WRITE (1, 190) Q, SPG, P21, P22, DELPG, T, CVG FORMAT (5X,'GAS FLOW RATE, ft^3/hr.:', T60, F9.3,/, 5X,'SPECIFIC GRAVITY:', T60, F9.3,/, 5X,'INLET PRESSURE, psia:', T60, F9.3,/, 5X,'OUTLET PRESSURE, psia:', T60, F9.3,/, 5X,'OUTLET PRESSURE, psia:', T60, F9.3,/, 190 * * 5X,'CONTROL VALVE PRESSURE DROP, psi:', T60, F9.3,/, 5X,'INLET TEMPERATURE, OF:', T60, F9.3,/, 5X,'CONTROL VALVE CAPACITY COEFFICIENT:', T60, F9.3) * ٠ . ENDIF WRITE (1, 200) 200 FORMAT (76(1H-)) GO TO 5 WRITE (1, 210) 66 FORMAT (//,20X, 'CONTROL VALVE SIZING FOR STEAM SERVICE'. 210 /,76(1H*)) с CALCULATE THE CONTROL VALVE FOR STEAM SERVICE. CALL STEAM WRITE (1, 220) CHOICE 220 FORMAT (5X, 'FLUID SERVICE:', T60, A) IF (CVS .GT. 0.0) THEN WRITE (1, 230) W, F, P31, P32, DELPS, CVS FORMAT (5X,'CALCULATED STEAM FLOW RATE, lb/hr.:', T60, F9.3,/, 5X,'DEGREE OF SUPERHEAT, oF:', T60, F9.3,/, 5X,'OUTLET PRESSURE, psia:', T60, F9.3,/, 5X,'CONTROL VALVE PRESSURE DROP, psi:', T60, F9.3,/, 5X,'CONTROL VALVE CAPACITY COEFFICIENT:', T60, F9.3) 230 ٠ * WRITE (1, 240) 240 FORMAT (76(1H-))

с FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE. WRITE (1, *) CHAR(12) ELSE WRITE (1, 250) W, F, P31, P32, DELPS, CVS WATE (1, 230) W, F, F31, F32, DELFS, CV3 FORMAT (5x,'STEAM FLOW RATE, 1b/hr.:', T60, F9.3,/, 5x,'DEGREE OF SUPERHEAT, oF:', T60, F9.3,/, 5x,'OUTLET PRESSURE, psia:', T60, F9.3,/, 5x,'CONTROL VALVE PRESSURE DROP, psi:', T60, F9.3,/, 5x,'CONTROL VALVE CAPACITY COEFFICIENT:', T60, F9.3) 250 ٠ * * * ٠ WRITE (1, 260) FORMAT (76(1H-)) 260 ENDIF с FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE. WRITE (1, *)CHAR(12)CLOSE(3,STATUS='KEEP') CLOSE (1) 999 STOP END с с с THIS PROGRAM CALCULATES THE CONTROL VALVE CAPACITY COEFFICIENT OR THE MAXIMUM FLOWRATE FOR FULLY OPEN CONTROL VALVE AND LIQUID с FLOW CONDITIONS. ****** ****** с SUBROUTINE LIQUID COMMON/DATA1/ V, P1, P2, DELPL, SPL, CVL CHECK WHETHER COMPUTATION REQUIRES CAPACITY COEFFICIENT OR с с MAXIMUM LIQUID FLOW RATE. с CALCULATE THE PRESSURE DROP ACROSS THE CONTROL VALVE, psi. DELPL = P1 - P2IF (CVL .GT. 0.0) THEN GO TO 10 ELSE CALCULATE THE VALVE CAPACITY COEFFICIENT (Cv) с CVL = V*SQRT(SPL)/SQRT(DELPL) ENDIF GO TO 30

399

- с CALCULATE THE MAXIMUM LIQUID FLOW RATE, qpm. V = CVL*SQRT(DELPL)/SQRT(SPL) 10 RETURN 30 END С THIS PROGRAM CALCULATES THE CONTROL VALVE CAPACITY COEFFICIENT OR с с с THE MAXIMUM GAS FLOW RATE FOR GAS FLOW CONDITIONS ¢ SUBROUTINE GAS COMMON/DATA2/ Q, P21, P22, DELPG, SPG, T, CVG T = T + 460.0с CALCULATE THE CONTROL VALVE PRESSURE DROP, psi. DELPG = P21 - P22 $VALG = P22 \star DELPG$ с с CHECK WHETHER COMPUTATION REQUIRES CAPACITY COEFFICIENT OR MAXIMUM GAS FLOW RATE. IF (CVG .GT. 0.0) THEN GO TO 10 ELSE ENDIF с CHECK WHETHER FLOW IS CRITICAL OR SUBCRITICAL IF ((P21/2.) .GE. P22) THEN GO TO 15 ELSE С CALCULATE THE VALVE CAPACITY COEFFICIENT FOR SUBCRITICAL FLOW CVG = Q*SQRT(SPG*T)/(1360.0*SQRT(VALG)) ENDIF GO TO 20 c c CALCULATE THE VALVE CAPACITY COEFFICIENT FOR CRITICAL FLOW. 15 WRITE (*, 25) 25 FORMAT (3X, 'CRITICAL GAS FLOW') pause CVG = Q*SQRT(SPG*T)/(1360.0*0.5*P21) GO TO 20 CALCULATE THE MAXIMUM GAS FLOW RATE, ft^3/hr. FOR SUBCRITICAL c c FLOW.
- 10 IF ((P21/2.0) .GE. P22) THEN

```
GO TO 35
       ELSE
        Q = (CVG*1360.0*SQRT(VALG))/SQRT(SPG*T)
        ENDIF
       GO TO 20
       CALCULATE THE MAXIMUM GAS FLOW RATE, ft^/hr. FOR CRITICAL FLOW
С
       WRITE (*, 45)
FORMAT(3X,'CRITICAL GAS FLOW')
 35
 45
       pause
       Q = (CVG*1360.0*0.5*P21)/SQRT(SPG*T)
20
       T = T - 460.0
       RETURN
       END
С
С
        *****
                                                                    ****
c
c
        THIS PROGRAM CALCULATES THE CONTROL VALVE CAPACITY COEFFICIENT
       OR THE MAXIMUM STEAM FLOW RATE FOR FULLY OPEN CONTROL VALVE AND
c
       FLOW CONDITIONS.
        SUBROUTINE STEAM
       REAL K
       COMMON/DATA3/ W, P31, P32, DELPS, F, CVS
       K = 1.0 + .0007 * F
с
       CALCULATE THE CONTROL VALVE PRESSURE DROP, psi.
       DELPS = P31 - P32
       VALS = P32*DELPS
c
c
       CHECK WHETHER COMPUTATION REQUIRES CAPACITY COEFFICIENT OR
       MAXIMUM FLOW RATE.
       IF (CVS .GT. 0.0) THEN
GO TO 10
        ELSE
        ENDIF
       CHECK WHETHER FLOW IS CRITICAL OR SUBCRITICAL
С
       IF ((P31/2.) .GE. P32) THEN
GO TO 15
       ELSE
       ENDIF
с
       CALCULATE THE CAPACITY COEFFICIENT FOR SUBCRITICAL FLOW
       CVS = (W*K)/3.*SQRT(VALS)
```

	GO TO 20
15 25	WRITE(*,25) Format(3X,'CRITICAL STEAM FLOW')
	pause
с	CALCULATE THE CAPACITY COEFFICIENT FOR CRITICAL FLOW.
	CVS = (W*K)/(1.5*P31)
	GO TO 20
c c	CHECK WHETHER FLOW IS CRITICAL OR SUBCRITICAL FLOW AND CALCULATE THE FLOW.
10	IF ((P31/2.) .GE. P32) THEN GO TO 35
	ELSE
с	CALCULATE THE MAXIMUM FLOW RATE FOR CRITICAL FLOW.
	W = 3.0*CVS*SQRT(VALS)/K ENDIF
	GO TO 20
35 45	WRITE(*, 45) Format(3x,'Critical steam flow')
	pause
с	CALCULATE THE MAXIMUM FLOW RATE FOR CRITICAL FLOW.
	₩=(1.5*P31*CVS)/K
20	RETURN END

PROGRAM PROG53

0000000		RELIEF-VALVE SIZING FOR VAPOR, STEAM, LIQUID, AIR AND FIRE. THE PROGRAM USES THE EQUATIONS AND CORRELATIONS PRESENTED IN AMERICAN PETROLEUM INSTITUTE (API) 520 RECOMMENDED PRACTICES FOR THE DESIGN AND INSTALLATION OF PRESSURE-RELIEVING SYSTEMS IN REFERINERIES FOR THE SIZING OF RELIEF VALVES IN VAPOR, STEAM, LIQUID, AIR AND FIRE SERVICES.
		CHARACTER COUNT(14),REGIME(14),REG(14),REG1(14) CHARACTER*14 COUNT1(14),ACOUNT(14),COUNLL(14),ACONT(14) CHARACTER*10 CHOICE
		REAL OP1, MW1, N1 REAL OP2, KSH, MW2, N2 REAL KV, KV1, KW, KD REAL OP4 REAL LAT, OP5, MW5, N5
		DIMENSION Y(14) COMMON/NAME/I,Y,IJK COMMON/DATA/REGI,ACONT COMMON/DATA/REGI,ACONT COMMON/DATA2/WSMAX,ADES1,ACAL1,C1 COMMON/DATA2/WSMAX,ADES2,ACAL2,C2 COMMON/DATA3/CPIMAX,GP2MAX,ADES3,ACAL3,ACAL31,ADES31,RE COMMON/DATA3/CPIMAX,GP2MAX,ADES3,ACAL3,ACAL31,ADES31,RE COMMON/DATA5/FORCE COMMON/DATA6/ADES4,VAMAX,ACAL4,C4 COMMON/DATA6/ADES4,VAMAX,ACAL4,C4 COMMON/DATA7/Q,WF COMMON/DATA8/KV1 COMMON CHOICE
	*	DATA COUNT/'D','E','F','G','H','J','K','L','M','N', 'P','Q','R','T'/
	* * * * * *	DATA COUNT1/' 1-2 ',' 1-2 ', ' 1.5-2 ',' 1.5-2 ', ' 1.5-3 or 2-3 ',' 2-3 or 2.5-4', ' 3-4 ',' 3-4 or 4-6 ', ' 4-6 ',' 4-6 ', ' 6-8 or 6-10 ',' 8-10 '/
		OPEN (UNIT=3, FILE='DATA53.DAT', STATUS='OLD', ERR=18) OPEN (UNIT =1, FILE ='PRN')
5		READ(3, 5, ERR=19) CHOICE FORMAT (A)
		<pre>IF (CHOICE .EQ. 'VAPOR' .OR. CHOICE .EQ. 'vapor') THEN GO TO 10 ELSEIF (CHOICE .EQ. 'STEAM' .OR. CHOICE .EQ. 'steam') THEN GO TO 20 ELSEIF (CHOICE .EQ. 'LIQUID' .OR. CHOICE .EQ. 'liquid') THEN GO TO 30 ELSEIF (CHOICE .EQ. 'AIR' .OR. CHOICE .EQ. 'air') THEN GO TO 40 ELSEIF (CHOICE .EQ. 'FIRE' .OR. CHOICE .EQ. 'fire') THEN</pre>

GO TO 50

ENDIF

c c	CALCULATE RELIEF VALVE FOR VAPOR CONDITION INPUT DATA FOR VAPOR CONDITION
0000000	<pre>WG = VAPOR FLOW RATE, lb/hr. SP1 = RELIEF VALVE SET PRESSURE, psig. OP1 = ALLOWABLE OVERPRESSURE, (%). T1 = RELIEVING TEMPERATURE OF THE INLET VAPOR (oF). Z1 = COMPRESSIBILITY FACTOR. MW1 = MOLECULAR WEIGHT OF VAPOR. N1 = RATIO OF THE SPECIFIC HEATS.</pre>
10	READ (3, *, ERR=19) WG, SP1 READ (3, *, ERR=19) OP1, T1 READ (3, *, ERR=19) Z1, MW1 READ (3, *, ERR=19) N1
с	DETERMINE THE RELIEF VALVE ORIFICE AREA FOR VAPOR CONDITION
	CALL VAPOR (WG, SP1, OP1, T1, Z1, MW1, N1)
С	PRINT RESULTS
	CALL OUT1 (WG, SP1, OP1, T1, Z1, NW1, N1)
	GO TO 60
с	INPUT DATA FOR STEAM RELIEF CONDITION.
0000000	WS=STEAM FLOW RATE, lb/hr.SP2=RELIEF VALVE SET PRESSURE, psig.OP2=ALLOWABLE OVERPRESSURE (%).KSH=SUPERHEAT STEAM CORRECTION FACTOR.T2=RELIEF TEMPERATURE, oF.MW2=MOLECULAR WEIGHT OF STEAM.N2=RATIO OF THE SPECIFIC HEATS.
20	READ (3, *, ERR=19) WS, SP2 READ (3, *, ERR=19) OP2, KSH READ (3, *, ERR=19) T2, MW2 READ (3, *, ERR=19) N2
С	DETERMINE THE RELIEF VALVE ORIFICE AREA FOR STEAM CONDITION
	CALL STEAM (WS, SP2, OP2, KSH, T2, MW2, N2)
с	PRINT RESULTS
	CALL OUT2 (WS, SP2, OP2, KSH, T2, MW2, N2)
	GO TO 60
с	INPUT DATA FOR LIQUID RELIEF
с с	GPM = LIQUID FLOW RATE, U.S. gpm. SP3 = UPSTREAM RELIEVING SET PRESSURE, psig. PB = TOTAL BACK PRESSURE, psig.

00000	SPL=SPECIFIC GRAVITY OF THE LIQUID.VISL=VISCOSITY OF LIQUID, CP.KV=CORRECTION FACTOR DUE TO VISCOSITY.KW=CORRECTION FACTOR DUE TO BACK PRESSURE.KD=EFFECTIVE COEFFICIENT OF DISCHARGE.
30	READ (3, *, ERR =19) GPM, SP3 READ (3, *, ERR = 19) PB, SPL READ (3, *, ERR=19) VISL, KV READ (3, *, ERR=19) KW, KD
с	DETERMINE THE RELIEF VALVE ORIFICE AREA FOR LIQUID CONDITION
	CALL LIQUID (GPM, SP3, PB, SPL, VISL, KV, KW, KD)
с	PRINT RESULTS
	IF (VISL .LT. 10.0) THEN
	CALL OUT3 (GPM, SP3, PB, SPL, VISL, KV, KW, KD)
	GO TO 60 ELSE
	CALL OUT32 (VISL)
	GO TO 60
	ENDIF
с	INPUT DATA FOR AIR RELIEF
с с с	VA = REQUIRED AIR CAPACITY, ft^3/min. T = INLET TEMPERATURE, oF. P = RELIEVING PRESSURE, psia. SP4 = RELIEF VALVE SET PRESSURE, psig.
40	READ (3, *, ERR=19) VA, T READ (3, *, ERR=19) P, SP4
с	DETERMINE THE RELIEF VALVE ORIFICE AREA FOR AIR CONDITION
	CALL AIR (VA, T, P, SP4)
с	PRINT RESULTS
	CALL OUT4 (VA, T, P, SP4)
	GO TO 60
с	DETERMINE THE RELIEF VALVE ORIFICE AREA FOR FIRE CONDITION.
00000000	F=ENVIRONMENTAL FACTOR.A=TOTAL WETTED SURFACE AREA, ft^2.LAT=LATENT HEAT OF VAPORIZATION, Btu/lb.SP5=RELIEF VALVE SET PRESSURE, psig.OP5=ALLOWABLE OVERPRESSURE, (%).T=RELIEVING TEMPERATURE, oF.Z=COMPRESSIBILITY FACTOR.MW5=FLUID MOLECULAR WEIGHT.

С	N5 = RATIO OF THE SPECIFIC HEATS.
с	INPUT DATA FOR FIRE RELIEF CONDITION.
50	READ (3, *, ERR=19) F, A READ (3, *, ERR=19) LAT, SP5 READ (3, *, ERR=19) OP5, T READ (3, *, ERR=19) Z, MW5 READ (3, *, ERR=19) N5
с	DETERMINE THE RELIEF VALVE ORIFICE AREA FOR FIRE CONDITION
	CALL FIRE (F, A, LAT, SP5, OP5, T, Z, MW5, N5)
с	PRINT RESULTS
	CALL OUT5 (F, A, LAT, SP5, OP5, T, Z, MW5, N5)
	GO TO 60
18 750	WRITE (*, 750) FORMAT (3X, 'DATA FILE DOES NOT EXIST')
	GO TO 60
19 800	WRITE (*, 800) FORMAT (3X, 'ERROR MESSAGE IN THE DATA VALUE')
	GO TO 60
	CLOSE (3, STATUS = 'KEEP') CLOSE (1)
60	STOP END
с с с	**************************************
	SUBROUTINE VAPOR (WG, SP1, OP1, T1, Z1, MW1, N1)
	CHARACTER COUNT(14), REGIME(14), REG(14) CHARACTER*14 COUNT1(14), COUNLL(14), ACOUNT(14) REAL K,KB,MMI,N1,OP1 COMMON/NAME/I,Y,IJK COMMON/DATA1/WGMAX,ADES1,ACAL1,C1 COMMON/DATA4/COUNT,REGIME,COUNT1,COUNLL,REG, ACOUNT COMMON/DATA5/FORCE DIMENSION Y(14)
с	CALCULATE THE GAS COEFFICIENT (C1) AND THE ORIFICE AREA.
	<pre>K=0.975 KB=1.0 T1=T1+460.0 P1=SP1+(SP1*OP1/100.0)+14.7 VAL1=N1*(2.0/(N1+1.0))**((N1+1.0)/(N1-1.0)) C1=520.0*SQRT(VAL1) VAL2=C1*K*P1*KB*SQRT(MW1)</pre>

ACAL1=WG*SORT(T1*Z1)/VAL2 DO 22 I=1, 14 IF (Y(I) .GT. ACAL1) THEN GO TO 25 ELSE ENDIE 22 CONTINUE ADES1=Y(I) 25 CALCULATE THE MAXIMUM FLOWRATE FROM THE MANUFACTURER'S С C ORIFICE AREA CALL MANUF(ADES1) WGMAX=ADES1*VAL2/SQRT(T1*21) CALCULATE THE REACTIVE FORCE С CALL REACTF(WGMAX,N1,T1,MW1) T1 = T1 - 460.0RETURN END С С THIS PROGRAM SIZES THE RELIEF-VALVE FOR STEAM CONDITION с ********** SUBROUTINE STEAM (WS, SP2, OP2, KSH, T2, MW2, N2) CHARACTER COUNT(14), REGIME(14), REG (14) CHARACTER*14 COUNT1(14), COUNLL(14), ACOUNT(14) COMMON/NAME/I,Y,IJK COMMON/DATA2/WSMAX, ADES2, ACAL2, C2 COMMON/DATA4/COUNT, REGIME, COUNT1, COUNLL, REG, ACOUNT DIMENSION Y(14) REAL KSH, OP2, MW2, N2, KN, KD KD = THE EFFECTIVE COEFFICIENT OF DISCHARGE (KD=0.975) С с KN = CORRECTION FACTOR FOR NAPIER EQUATION (KN=1) с CALCULATE THE GAS COEFFICIENT (C2) AND THE ORIFICE AREA KN = 1.0KD = 0.975T2 =T2+460.0 P2=SP2+(SP2*OP2/100.0)+14.7 VAL3=N2*(2.0/(N2+1.0))**((N2+1.0)/(N2-1.0)) C2=520.0*SQRT(VAL3) ACAL2=WS/(51.5*P2*KSH*KD*KN) DO 33 I=1,14 IF (Y(I) .GT. ACAL2) THEN GO TO 25 ELSE ENDIF 33 CONTINUE CALCULATE THE MAXIMUM FLOWRATE FROM THE MANUFACTURER'S ORIFICE С с AREA.

```
ADES2=Y(I)
 25
       CALL MANUF(ADES2)
       WSMAX=51.5*P2*KSH*KN*KD*ADES2
С
       CALCULATE THE REACTIVE FORCE.
       CALL REACTF(WSMAX,N2,T2,MW2)
       T2 = T2 - 460.0
       RETURN
       END
       ***********
С
       THIS PROGRAM SIZES THE RELIEF VALVE FOR LIQUID CONDITION.
С
       SIZING FOR LIQUID RELIEF VALVES REQUIRING LIQUID CAPACITY
c
С
       CERTIFICATION.
        *****
с
       SUBROUTINE LIQUID (GPM, SP3, PB, SPL, VISL, KV, KW, KD)
       DIMENSION Y(14)
       CHARACTER COUNT(14), REGIME(14), REG(14), REG1(14)
       CHARACTER*14 COUNT1(14), ACOUNT(14), COUNLL(14), ACONT(14)
       REAL KD, KW, KV, KV1
       COMMON/DATA3/GP1MAX, GP2MAX, ADES3, ACAL3, ACAL31, ADES31, RE
       COMMON/NAME/I,Y, IJK
       COMMON/DATA4/COUNT, REGIME, COUNT1, COUNLL, REG, ACOUNT
       COMMON/DATA8/KV1
       COMMON/DATA/REG1, ACONT
С
       CALCULATE THE ORIFICE AREA FOR LIQUID CAPACITY CERTIFICATION.
       SP3 = SP3+(0.1*SP3)
       VAL3 = (SP3-PB)
       VAL4=38.0*KD*KW*KV*SQRT(VAL3)
       ACAL3=GPM*SQRT(SPL)/VAL4
       DO 44 I =1, 14
       IF (Y(I) .GT. ACAL3) THEN
           GO TO 25
           ELSE
       ENDIF
44
       CONTINUE
25
       ADES3=Y(I)
       IJK = I
с
       CALCULATE THE MANUFACTURER'S ORIFICE AREA AND THE MAXIMUM
С
С
       LIQUID FLOWRATE.
       CALL MANUF(ADES3)
       GP1MAX=(ADES3*VAL4)/SQRT(SPL)
       CALL OUT3 (GPM, SP3, PB, SPL, VISL, KV, KW, KD)
```

IF (VISL .GE. 10.) THEN GO TO 30 ELSE ENDIF GO TO 90 30 CALL AMANU(ADES3) REG1(I)=REG(I) ACONT(I)=ACOUNT(I) GP1MAX=(ADES3*VAL4)/SQRT(SPL) CALL OUT31(GPM, SP3, PB, SPL, VISL, KV, KW, KP) с RE=(GPM*2800.0*SPL)/(VISL*SQRT(ADES3)) VAL6=(1.2502*ALOG(RE))+(0.17510*(ALOG(RE))**2) VAL7=(0.01087*(ALOG(RE))**3)-(0.00025*(ALOG(RE))**4) KV1=-2.38878+VAL6+VAL7 VAL8=38.0*KD*KW*KV1*SQRT(VAL3) ACAL31=GPM*SQRT(SPL)/VAL8 С DO 66 I=1,14 IF (Y(I) .GT. ACAL31) THEN GO TO 77 ELSE ENDIF CONTINUE 66 77 ADES31=Y(I) CALL MANUFL (ADES3, ADES31) GP2MAX=(ADES31*VAL8)/SQRT(SPL) 90 RETURN END С С THIS PROGRAM PRINTS THE RESULTS OF VAPOR RELIEF SIZING ONTO A С PRINTER С ************ SUBROUTINE OUT1 (WG, SP1, OP1, T1, Z1, MW1, N1) CHARACTER COUNT(14),REGIME(14),REG(14),REG1(14) CHARACTER*14 COUNT1(14),ACOUNT(14),COUNLL(14),ACONT(14) CHARACTER*10 CHOICE COMMON/NAME/I,Y,IJK COMMON/DATA1/WGMAX, ADES1, ACAL1, C1 COMMON/DATA4/COUNT, REGIME, COUNT1, COUNLL, REG, ACOUNT COMMON/DATA5/FORCE COMMON CHOICE REAL OP1, MW1, N1 DIMENSION Y(14)

WRITE(1, 100)
100 FORMAT(20X,'VAPOR RELIEF VALVE SIZING')
WRITE(1, 110)

105 110 120	WRITE (1, 105)CHOICE FORMAT (3X,'BASIS OF SELECTION:', T65, A) FORMAT(1H ,76(1H*)) WRITE(1, 120)WG FORMAT(3X,'REQUIRED VAPOR RATE, lb/hr.:',T65, F12.0)
130 140	WRITE(1, 130) SP1 FORMAT(3X,'VALVE SET PRESSURE, psig:',T65, F9.3) WRITE(1, 140) OP1 FORMAT(3X,'PERCENTAGE IN OVERPRESSURE, (%):',T65, F9.3)
150 160	WRITE(1, 150)T1 FORMAT(3X,'UPSTREAM FLUID TEMPERATURE, DEG. F:',T65, F9.3) WRITE(1, 160)Z1 FORMAT(3X,'COMPRESSIBILITY FACTOR:',T65, F9.4)
170	WRITE(1, 170) MW1 FORMAT(3X,'MOLECULAR WEIGHT OF VAPOR:', T65, F9.3)
180	WRITE(1, 180)N1 FORMAT(3X,'RATIO OF SPECIFIC HEATS OF FLUID:', T65, F9.3)
190	WRITE(1, 190) C1 FORMAT(3X,'GAS CONSTANT DETERMINED BY RATIO OF Cp/Cv:',T65,F9.3)
200	WRITE(1, 200) ACAL1 FORMAT(3X,'CALCULATED VALVE ORIFICE AREA, in^2:',T65, F9.3)
210	WRITE(1, 210) ADES1 FORMAT(3X,'NEAREST STANDARD ORIFICE AREA, in^2:',T65, F9.3)
220	WRITE(1, 220) REGIME(I) FORMAT(3X,'NEAREST STANDARD ORIFICE SIZE:', T65, A)
230	WRITE(1, 230) COUNT1 (I) FORMAT(3X,'PREFERRED VALVE BODY SIZE: (IN-OUT) inches:',T65,A14)
240 *	WRITE(1, 240) WGMAX FORMAT(3X,'MAXIMUM VAPOR FLOWRATE WITH STD. ORIFICE, lb/hr.:', T65,F12.0)
250	WRITE(1, 250) FORCE FORMAT(3X,'REACTION FORCE, 1b:',T65, F9.0)
260	WRITE (1, 260) FORMAT (76(1H-))
с	FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.
	WRITE (1, *)CHAR(12)
20	RETURN STOP END
c c	**************************************

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c c	ONTO A PRINTER ************************************
	SUBROUTINE OUT3 (GPM, SP3, PB, SPL, VISL, KV, KW, KD) DIMENSION Y(14) CHARACTER COUNT(14), REGINE(14), REG(14), REG1(14) CHARACTER*14 COUNT1(14), ACOUNT(14), COUNLL(14), ACONT(14) CHARACTER*10 CHOICE COMMON/NAME/I, Y, JJX COMMON/DATA/REG1, ACONT COMMON/DATA/REG1, ACONT COMMON/DATA/GPIMAX, GP2MAX, ADES3, ACAL3, ACAL31, ADES31, RE COMMON/DATA/COUNT, REGIME, COUNT1, COUNLL, REG, ACOUNT COMMON/DATA/KV1 COMMON CHOICE REAL KV, KW, KD, KV1
270	WRITE(1, 270) FORMAT(20X,'RELIEF VALVE SIZING FOR LIQUID') WRITE(1, 280)
280	FORMAT(1H ,76(1H*))
285	WRITE (1, 285)CHOICE FORMAT (3X,'BASIS OF SELECTION:', T65, A) WRITE(1, 290) GPM
290	FORMAT(3X,'LIQUID FLOWRATE, US gpm:',T65, F12.0) WRITE(1, 300) SP3
300	<pre>FORMAT(3X,'VALVE SET PRESSURE, psig:',T65, F9.3) WRITE(1,310) PB</pre>
310 320	FORMAT(3X,'BACK PRESSURE ON VALVE, psig:',T65, F9.3) WRITE(1, 320) SPL FORMAT(3X,'FLUID SPECIFIC GRAVITY:', T65, F9.3)
330	WRITE(1,330) VISL FORMAT(3X,'FLUID VISCOSITY, cP:', T65, F9.4)
340	WRITE(1, 340) KV FORMAT(3X,'VISCOSITY CORRECTION FACTOR, Kv:',T65, F9.4)
350	WRITE(1, 350) KW FORMAT(3X,'BACKPRESSURE CORRECTION FACTOR, Kw:',T65, F9.4)
360	<pre>WRITE(1, 360) KD FORMAT(3X,'EFFECTIVE COEFFICIENT OF DISCHARGE, Kd:', T65, F9.3) WRITE(1, 370) ACAL3</pre>
370	FORMAT(3X,'CALCULATED VALVE ORIFICE AREA, in^2:',T65,F9.3)
380	WRITE(1, 380) ADES3 FORMAT(3X,'NEAREST STANDARD ORIFICE AREA, in^2:', T65,F9.3)
390	WRITE(1, 390) REGIME (I) FORMAT(3X,'NEAREST STANDARD ORIFICE SIZE:', T65, A)
400	<pre>WRITE(1, 400) COUNLL(I) FORMAT(3X,'PREFERRED VALVE BODY SIZE, (IN-OUT) inches:',T65,A14)</pre>
410 *	<pre>WRITE(1, 410) GP1MAX FORMAT(3X,'MAXIMUM LIQUID FLOWRATE WITH STD. ORIFICE, gpm:',</pre>

FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE. C RETURN END с ********* SUBROUTINE OUT31 (GPM, SP3, PB, SPL, VISL, KV, KW, KD) CHARACTER COUNT(14), REGIME(14), REG(14), REG1(14) CHARACTER*14 COUNT1(14), ACOUNT(14), COUNLL(14), ACONT(14) CHARACTER*10 CHOICE DIMENSION Y(14) COMMON/NAME/I,Y,IJK COMMON/DATA/REG1, ACONT COMMON/DATA3/GP1MAX, GP2MAX, ADES3, ACAL3, ACAL31, ADES31, RE COMMON/DATA4/COUNT, REGIME, COUNT1, COUNLL, REG, ACOUNT COMMON/DATA8/KV1 COMMON CHOICE REAL KV, KW, KD, KV1 WRITE(1, 420) FORMAT(20X, 'RELIEF VALVE SIZING FOR LIQUID') 420 WRITE(1, 430) FORMAT(1H ,76(1H*)) 430 WRITE (1, 440)CHOICE FORMAT (3X, 'BASIS OF SELECTION:', T65, A) 440 WRITE(1, 450) GPM 450 FORMAT(3X,'LIQUID FLOW RATE, U.S gpm:',T65,F12.0) WRITE(1, 460) SP3 FORMAT(3X, 'VALVE SET PRESSURE, psig:', T65, F9.3) 460 WRITE(1,470) SPL 470 FORMAT(3X, 'FLUID SPECIFIC GRAVITY:', T65, F9.3) WRITE(1, 480) KV 480 FORMAT(3X, 'VISCOSITY CORRECTION FACTOR Kv=1.0:', T65, F9.3) WRITE(1, 490) KW 490 FORMAT(3X, 'BACK PRESSURE CORRECTION FACTOR Kw=1.0:', T65, F9.3) WRITE(1, 500)KD 500 FORMAT(3X, 'COEFFICIENT OF DISCHARGE Kd=0.65:', T65, F9.3) WRITE(1, 510) ACAL3 FORMAT(3X, 'CALCULATED VALVE ORIFICE AREA, in^2:', T65, F9.3) 510 WRITE(1, 520) ADES3 520 FORMAT(3X, 'NEAREST STANDARD ORIFICE AREA, in^2:', T65, F9.3) WRITE(1, 530) REG1(I) FORMAT(3X, 'NEAREST STANDARD ORIFICE SIZE:', T65, A) 530 WRITE(1, 540) ACONT(I) FORMAT(3X, 'PREFERRED VALVE BODY SIZE, (IN-OUT) inches:', 540 T65,A14) WRITE(1, 550) GP1MAX FORMAT(3X, 'MAXIMUM LIQUID FLOWRATE WITH STD. ORIFICE, gpm:', 550 T65.F12.0) WRITE(1, 560)

560	FORMAT (76(1H-))
с	FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.
-	WRITE (1, *)CHAR(12)
	RETURN END
с с с	**************************************
	SUBROUTINE OUT32 (VISL) DIMENSION Y(14) CHARACTER COUNT(14),REGIME(14),REG(14),REG1(14) CHARACTER*14 COUNT1(14),ACOUNT(14),COUNLL(14),ACONT(14) COMMON/NAME/I,Y, IJK COMMON/DATA3/GPIMAX,GP2MAX,ADES3,ACAL3,ACAL31,ADES31,RE COMMON/DATA3/COUNT,REGIME,COUNT1,COUNLL, REG, ACOUNT COMMON/DATA8/KV1 REAL KV1
570	WRITE(1, 570) KV1 FORMAT(3X,'VISCOSITY CORRECTION FACTOR:', T65, F9.3)
580	WRITE(1, 500) RE FORMAT(3X,'REYNOLDS NUMBER:', T65, F12.0)
590	WRITE(1, 590) ACAL31 FORMAT(3X,'CALCULATED VALVE ORIFICE AREA, in^2:',T65,F9.3)
600	WRITE(1, 600) ADES31 FORMAT(3X,'NEAREST STANDARD ORIFICE AREA, in^2:', T65,F9.3)
610	WRITE(1, 610) REGIME (I) FORMAT(3X,'NEAREST STANDARD ORIFICE SIZE:', T65, A)
620	<pre>WRITE(1, 620) COUNLL (I) FORMAT(3X,'PREFERRED VALVE BODY SIZE, (IN-OUT) inches:',T65,A14)</pre>
630 *	<pre>WRITE(1, 630) GP2MAX FORMAT(3X,'MAXIMUM LIQUID FLOWRATE WITH STD. ORIFICE, gpm:',</pre>
640	WRITE (1, 640) FORMAT (76(1H-))
с	FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.
	WRITE (1, *)CHAR(12)
	RETURN END

```
С
        *****
С
        THIS PROGRAM PRINTS THE RESULTS OF STEAM RELIEF SIZING
С
        ONTO A PRINTER
С
        *****
        SUBROUTINE OUT2 (WS, SP2, OP2, KSH, T2, MW2, N2)
        CHARACTER COUNT(14), REGIME(14), REG(14)
        CHARACTER*14 COUNT1(14), COUNLL(14), ACOUNT(14)
        CHARACTER*10 CHOICE
        COMMON/NAME/I,Y,IJK
        COMMON/DATA2/WSMAX, ADES2, ACAL2, C2
        COMMON/DATA4/COUNT, REGIME, COUNT1, COUNLL, REG, ACOUNT
        COMMON/DATA5/FORCE
        COMMON CHOICE
        DIMENSION Y(14)
        REAL KSH, MW2, N2
        WRITE(1, 650)
 650
        FORMAT(20X, 'RELIEF VALVE SIZING FOR STEAM CONDITION')
        WRITE(1, 660)
        FORMAT(1H ,76(1H*))
 660
        WRITE (1, 665)CHOICE
 665
        FORMAT (3X, 'BASIS OF SELECTION:', T65, A)
        WRITE(1, 670) WS
        FORMAT(3X, 'REQUIRED STEAM RATE, 1b/hr.:', T65, F12.0)
 670
        WRITE(1, 680)SP2
 680
        FORMAT(3X, 'VALVE SET PRESSURE, psig:', T65, F9.3)
        WRITE(1, 690) OP2
 690
        FORMAT(3X, 'PERCENTAGE OF OVERPRESSURE, (%):', T65, F9.3)
        WRITE(1, 700) KSH
 700
        FORMAT(3X, 'FACTOR FOR SUPERHEATED KSH=0.88/SATURATED KSH=1:',
               T65,F9.3)
        WRITE(1, 710) T2
 710
        FORMAT(3X, 'STEAM TEMPERATURE, DEG. F:', T65, F9.3)
        WRITE(1, 720) MW2
 720
        FORMAT(3X, 'MOLECULAR WEIGHT:', T65, F9.3)
        WRITE(1,
                730) N2
        FORMAT(3X, 'RATIO OF SPECIFIC HEATS OF STEAM:', T65, F9.3)
 730
        WRITE(1, 740) C2
 740
        FORMAT(3X,'GAS CONSTANT DETERMINED BY RATIO OF Cp/Cv:', T65, F9.3)
        WRITE(1, 750) ACAL2
 750
        FORMAT(3X, 'CALCULATED VALVE ORIFICE AREA, in^2:', T65, F9.3)
        WRITE(1,
                760) ADES2
 760
        FORMAT(3X,'NEAREST STANDARD AREA, in^2:', T65, F9.3)
        WRITE(1, 770) REGIME (I)
 770
        FORMAT(3X, 'NEAREST STANDARD ORIFICE SIZE:', T65, A)
        WRITE(1, 780) COUNT1(I)
 780
        FORMAT(3X, 'PREFERRED VALVE BODY SIZE, (IN-OUT) inches: ', T65, A14)
        WRITE(1, 790) WSMAX
 790
       FORMAT(3X, 'MAXIMUM STEAM FLOWRATE WITH STD. ORIFICE, 1b/hr.:',
               T65,F12.0)
       WRITE(1, 800)FORCE
 800
        FORMAT(3X, 'REACTION FORCE, 1b:', T65, F12.0)
       WRITE (1, 810)
810
       FORMAT (76(1H-))
С
       FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.
       WRITE (1, *)CHAR(12)
       RETURN
       END
```

```
С
       č
       THIS PROGRAM SIZES THE RELIEF VALVE FOR AIR-CONDITION
       ******************
С
       SUBROUTINE AIR (VA, T4, SP4, OP4)
       REAL OP4, N, MW, KB, KD
       CHARACTER COUNT(14), REGIME(14), REG(14)
       CHARACTER*14 COUNT1(14), COUNLL(14), ACOUNT(14)
       COMMON/NAME/I,Y,IJK
       COMMON/DATA4/COUNT, REGIME, COUNT1, COUNLL, REG, ACOUNT
       COMMON/DATA5/FORCE
       COMMON/DATA6/ADES4,VAMAX,ACAL4,C4
       DIMENSION Y(14)
С
       CALCULATE THE ORIFICE AREA AND THE GAS COEFFICIENT.
       KB =1.0
       KD=0.953
       T = T4 + 460.0
       MW=28.97
       N=1.40
       P4=SP4+(SP4*OP4/100.0)+14.7
       RHO=(MW*P4)/(10.72*T)
       ACAL4=(VA*SQRT(T))/(418.0*KB*KD*P4)
       VAL4=N*(2.0/(N+1.0))**((N+1.0)/(N-1.0))
       C4=520.0*SORT(VAL4)
       DO 44 I=1,14
       IF (Y (I) .GT. ACAL4) THEN
          GO TO 30
          ELSE
       ENDIF
44
       CONTINUE
C
       CALCULATE THE MANUFACTURER'S AREA AND MAXIMUM AIR RATE.
30
       ADES4=Y(I)
       CALL MANUF(ADES4)
       VAMAX=(418.0*ADES4*P4*KB*KD)/SQRT(T)
       WAMAX=(60.0*VAMAX*RHO)
с
       CALCULATE THE REACTIVE FORCE.
       FORCE = WAMAX/15.9
       RETURN
      END
С
       *****
c
       THIS PROGRAM PRINTS THE RESULTS OF AIR RELIEF VALVE SIZING ONTO
С
       A PRINTER
с
       SUBROUTINE OUT4 (VA, T4, SP4, OP4)
       CHARACTER*14 COUNT1(14), COUNLL(14), ACOUNT(14)
       CHARACTER*10 CHOICE
       CHARACTER COUNT(14), REGIME(14), REG (14)
       COMMON/NAME/I,Y,IJK
      COMMON/DATA4/COUNT, REGIME, COUNT1, COUNLL, REG, ACOUNT
       COMMON/DATA5/FORCE
      COMMON/DATA6/ADES4,VAMAX,ACAL4,C4
      COMMON CHOICE
```

```
DIMENSION Y(14)
       REAL OP4
       WRITE(1, 820)
        FORMAT(20X, 'RELIEF VALVE SIZE FOR AIR CONDITION')
 820
       WRITE(1, 830)
       FORMAT(1H ,76(1H*))
 830
       WRITE (1, 835)CHOICE
       FORMAT (3X, 'BASIS OF SELECTION:', T65, A)
 835
       WRITE(1, 840)VA
       FORMAT(3X, 'REQUIRED AIR FLOWRATE, S.ft^3/min:', T65, F12.0)
 840
       WRITE(1, 850) T4
        FORMAT(3X,'AIR TEMPERATURE, DEG. F:', T65, F9.3)
 850
       WRITE(1, 860) SP4
        FORMAT(3X, 'VALVE SET PRESSURE, psig:', T65, F9.3)
 860
       WRITE(1, 870) OP4
       FORMAT(3X, 'PERCENTAGE OF OVERPRESSURE, (%):', T65, F9.3)
 870
       WRITE(1, 880) C4
        FORMAT(3X, 'GAS CONSTANT DETERMINED BY RATIO OF Cp/Cv:', T65, F9.3)
 880
       WRITE(1, 890) ACAL4
       FORMAT(3X, 'CALCULATED VALVE ORIFICE AREA, in^2:', T65, F9.3)
 890
       WRITE(1, 900) ADES4
       FORMAT(3X, 'NEAREST STANDARD AREA, in^2:', T65, F9.3)
 900
       WRITE(1, 1000) REGIME (I)
        FORMAT(3X, 'NEAREST STANDARD ORIFICE SIZE: ', T65, A)
 1000
       WRITE(1, 1100) ACOUNT(I)
       FORMAT(3X, 'PREFERRED VALVE BODY SIZE, (IN-OUT) inches: ', T65, A14)
 1100
       WRITE(1, 1200) VAMAX
       FORMAT(3X, 'MAXIMUM AIR FLOWRATE WITH STD. ORIFICE, ft^3/min:',
 1200
              T65,F12.0)
       WRITE(1, 1300) FORCE
       FORMAT(3X, 'REACTION FORCE, 1b:', T65, F12.0)
1300
       WRITE (1, 1400)
       FORMAT (76(1H-))
1400
       FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.
С
       WRITE (1, *)CHAR(12)
       RETURN
       END
        *********
С
       THIS PROGRAM SIZES THE RELIEF VALVE FOR FIRE CONDITION
С
c
        ********
                                                                  ****
       SUBROUTINE FIRE (F, AREA, LAT, SP5, OP5, T5, Z5, MW5, N5)
       REAL LAT, OP5, MW5, N5
       CHARACTER COUNT(14), REGIME(14), REG (14)
       CHARACTER*14 COUNT1(14), COUNLL(14), ACOUNT(14)
       COMMON/NAME/I,Y,IJK
       COMMON/DATA1/WGMAX, ADES1, ACAL1, C1
       COMMON/DATA4/COUNT, REGIME, COUNT1, COUNLL, REG, ACOUNT
       COMMON/DATA5/FORCE
        COMMON/DATA7/Q,WF
       DIMENSION Y(14)
С
       CALCULATE THE TOTAL HEAT ABSORBED
       Q=21000.0*F*(AREA)**0.82
```

C CALCULATE THE VAPOR RATE

WF=Q/LAT С CALCULATE THE ORIFICE AREA FOR FIRE CONDITION. CALL VAPOR (WF, SP5, OP5, T5, Z5, MW5, N5) RETURN END *********** С THIS PROGRAM PRINTS THE RESULTS OF FIRE RELIEF SIZING ONTO A С С PRINTER С SUBROUTINE OUT5 (F, AREA, LAT, SP5, OP5, T5, Z5, NW5, N5) CHARACTER COUNT(14), REGIME(14), REG(14) CHARACTER*14 COUNT1(14), COUNLL(14), ACOUNT(14) CHARACTER*10 CHOICE COMMON/NAME/I,Y,IJK COMMON/DATA1/WGMAX, ADES1, ACAL1, C1 COMMON/DATA4/COUNT, REGIME, COUNT1, COUNLL, REG, ACOUNT COMMON/DATA5/FORCE COMMON/DATA7/Q,WF COMMON CHOICE DIMENSION Y(14) REAL LAT, OP5, MW5, N5 C WRITE(1, 1500) FORMAT(20X, 'FIRE RELIEF VALVE SIZING') 1500 WRITE(1, 1600) FORMAT(1H ,76(1H*)) 1600 WRITE (1, 1650)CHOICE 1650 FORMAT (3X, 'BASIS OF SELECTION:', T65, A) WRITE(1, 1700) F 1700 FORMAT(3X, 'ENVIRONMENTAL FACTOR: ', T65, F9.3) WRITE(1, 1800) AREA 1800 FORMAT(3X,'WETTED SURFACE AREA OF VESSEL, ft^2:', T65, F9.3) WRITE(1, 1900) LAT FORMAT(3X, 'LATENT HEAT OF VAPORIZATION, Btu/lb:', T65, F9.3) 1900 WRITE(1, 2000) SP5 2000 FORMAT(3X, 'VALVE SET PRESSURE, psig:', T65, F9.3) WRITE(1, 2100) OP5 2100 FORMAT(3X, 'PERCENTAGE OF OVERPRESSURE, (%):', T65, F9.3) WRITE(1, 2200) T5 2200 FORMAT(3X, 'FLUID TEMPERATURE, DEG. F:', T65, F9.3) WRITE(1, 2300) Z5 2300 FORMAT(3X, 'FLUID COMPRESSIBILITY FACTOR: ', T65, F9.3) WRITE(1, 2400) MW5 2400 FORMAT(3X, 'FLUID MOLECULAR WEIGHT:', T65, F9.3) WRITE(1, 2500) N5 2500 FORMAT(3X, 'RATIO OF SPECIFIC HEATS OF FLUID: ', T65, F9.3) WRITE(1, 2600) C1 FORMAT(3X, 'GAS CONSTANT DETERMINED BY RATIO OF Cp/Cv:', T65, F9.3) 2600 WRITE(1, 2700) WF 2700 FORMAT(3X, 'VAPOR RELIEF FLOW RATE, 1b/h.:', T65, F12.0) WRITE(1, 2800) ACAL1 2800 FORMAT(3X, 'CALCULATED VALVE ORIFICE AREA, in^2:', T65, F9.3) WRITE(1, 2900) ADES1 2900 FORMAT(3X, 'NEAREST STANDARD ORIFICE AREA, in^2:', T65, F9.3)

```
WRITE(1, 3000) REGIME(I)
       FORMAT(3X, 'NEAREST STANDARD ORIFICE SIZE:', T65, A)
3000
       WRITE(1, 3100) COUNT1(I)
       FORMAT(3X, 'PREFERRED VALVE BODY SIZE, (IN-OUT) inches: ', T65, A14)
3100
       WRITE(1, 3200) WGMAX
       FORMAT(3X, 'MAXIMUM VAPOR FLOWRATE WITH STD. ORIFICE lb/hr.:',
3200
             T65,F12.0)
       WRITE(1, 3300) FORCE
       FORMAT(3X, 'REACTION FORCE, 1b:', T65, F12.0)
3300
       WRITE (1, 3400)
       FORMAT (76(1H-))
3400
       FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.
С
       WRITE (1, *)CHAR(12)
       RETURN
       STOP
       END
       *************************
с
       THIS PROGRAM STORES THE MANUFACTURERS' ORIFICE SIZES
С
             С
       ***
       SUBROUTINE MANUF (ADES)
       CHARACTER COUNT(14), REGIME(14), REG(14)
       CHARACTER*14 COUNT1(14), ACOUNT(14), COUNLL(14)
       COMMON/NAME/I,Y, IJK
       COMMON/DATA4/COUNT, REGIME, COUNT1, COUNLL, REG, ACOUNT
       DIMENSION Y(14)
       DO 10 I≔1,14
       IF (ADES .EQ. Y(I)) THEN
              REGIME(I)=COUNT(I)
              COUNLL(I)=COUNT1(I)
              GO TO 20
       ELSE
       ENDIF
10
       CONTINUE
20
       RETURN
       END
с
С
       c
c
       THIS PROGRAM STORES THE MANUFACTURERS' ORIFICE SIZES
       SUBROUTINE AMANU (ADES3)
       CHARACTER COUNT(14), REGIME(14), REG(14), REG1(14)
       CHARACTER*14 COUNT1(14), ACOUNT(14), COUNLL(14), ACONT(14)
       COMMON/NAME/I,Y,IJK
       COMMON/DATA/REG1, ACONT
       COMMON/DATA4/COUNT, REGIME, COUNT1, COUNLL, REG, ACOUNT
       DIMENSION Y(14)
       DO 10 I=1,14
          IF (ADES3 .EQ. Y(I)) THEN
          REG(I)=COUNT(I)
          ACOUNT(I)=COUNT1(I)
          ELSE
       ENDIF
10
       CONTINUE
```

0 0 0 0	RETURN END THIS PROGRAM STORES THE MANUFACTURERS' ORIFICE SIZES SUBROUTINE MANUFL (ADES3, ADES31) CHARACTER COUNT(14),REGIME(14),REG(14) CHARACTER*14 COUNT(14),ACOUNT(14),COUNLL(14) COMMON/NAME/I,Y,IJK COMMON/NAME/I,Y,IJK COMMON/NATA4/COUNT,REGIME,COUNT1,COUNLL, REG, ACOUNT DIMENSION Y(14) DO 10 I=1, 14
10	IF (ADES3 .EQ. Y(I)) THEN REG(I)=COUNT(I) ACOUNT(I)=COUNT1(I) ELSE ENDIF CONTINUE
10	DO 20 I=1,14 IF (ADES31 .EQ. Y(I)) THEN REGIME(I)=COUNT(I) COUNLL(I)=COUNT1(I) GO TO 30 ELSE
20 30	ENDIF CONTINUE RETURN END
с с с с	<pre>this program determines the reaction forces in an open discharge system subroutine reactf (wg, n, t, Mw) real n, Mw common/Data5/Force Val1=(N*T)/((N+1.0)*Mw) Force=(wg*SQRT(Val1))/366.0 return END</pre>
с	<pre>Here the second se</pre>

CHAPTER 6

Compressors

INTRODUCTION

Compressors are gas movers and devices where mechanical work is done on the gas resulting in an increased pressure. Centrifugal compressors are used in a variety of chemical process industries (CPI). Their performance often varies with changes in process conditions. Sometimes the performance curves supplied by the manufacturer as discharge pressure and power requirement against an inlet volumetric flow may not be valid for variations in process conditions. Lapina [1] has provided a technique of obtaining a usable performance curve that is valid for the actual process conditions. Gas compressor applications are employed in the following duties:

- refrigeration
- gas injection
- gas transmission
- gas lift
- wellhead separator boosting

There is no single type of compressor that can be adapted to a particular application. The operating conditions, space, and weight restrictions must be reviewed before the appropriate compressor is selected. In addition, the type of driver requires selection because its selection and process conditions are interrelated.

Compressors can be classified into two basic types: reciprocating and centrifugal. The reciprocating compressor is well suited for high pressures and low volume flow rates, while the centrifugal is preferred for lower pressures and high volume flow rates. A centrifugal compressor uses the relationship between the velocity and pressure to increase the gas pressure. The gas enters a rotating impeller at the eye. The vanes force the gas to the outside rim, and subsequently throw it away from the rim at a high velocity. The gas is flung into the surrounding diffuser and volute passageways with a large volume, resulting in a reduced velocity. The velocity energy is thus changed into pressure energy and subsequently increases the gas pressure.

CENTRIFUGAL COMPRESSORS

In a centrifugal compressor, at least half the driver power (e.g., an electric gas or a steam turbine or a gas engine) is changed into gas velocity energy. The increased velocity can be carried forward to the next impeller for further increase in the velocity. A centrifugal compressor uses the impeller to furnish rotational pressure, which squeezes the gas outward and closer together, resulting in some pressure increase. As the impeller is rotated faster, more energy is expended from the driver. Figure 6-1 shows the flow of gas in a centrifugal compressor. The amount of pressure a compressor impeller can develop depends on its speed and diameter. Increased velocity and pressure are required for high speed, large diameter impellers. The impeller metal will have the same centrifugal force as the gas flowing through it. Also the impeller metal can tear apart if the diameter is too large or the impeller turns too rapidly. Therefore, the pressure that is developed with a single impeller is

CENTRIFUGAL FORCE

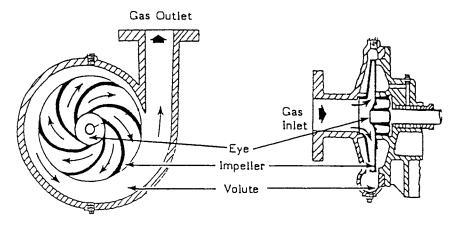


Figure 6-1. Gas flow in centrifugal compressor.

limited by the strength of the impeller metal. An additional impeller is used where one impeller is incapable of providing the required pressure. The gas leaving one rim enters the next stage eye of the impeller, consequently boosting its pressure.

RECIPROCATING COMPRESSORS

Reciprocating compressors are furnished as single stage or multistage with ratings from fractional to more than 20,000 horsepower per unit. The pressures range from a low vacuum at suction to 30,000 psi and higher at discharge. The number of stages depends on the overall compression ratio. The compression ratio per stage is limited by the discharge temperature and does not exceed four [2].

Multistage compressors are provided with intercoolers between stages and with an aftercooler at the compressor discharge. These are heat exchangers that remove the heat of compression from the gas and reduce its temperature to approximately the temperature existing at the compressor intake. In certain cases, aftercooling and interstage cooling are not always beneficial. If the gas must be heated before entering a reactor, downstream heaters can be designed for a lower duty if the heat of compression is not removed. Multistage compressors are intercooled when the inlet temperature of the gas and the required compression ratio are such that the discharge temperature of the gas exceeds 300°F. The coolers reduce the actual volume of gas flowing to the high pressure cylinders, the horsepower requirement, and keep the temperature within safe operating limits. In instances where a compressor with intercoolers is used to compress gases with condensable vapors, the combined effect of compression and cooling can condense out the liquid. Therefore, the liquid condensate must be removed by installing a knockout pot after the intercooler. This will prevent damage to the downstream of the compressor.

THE EQUATIONS

Compressors are rated in feet of compression head developed. This is the energy conveyed to a gas stream by a compressor. It is observed by the increase in gas pressure as the gas passes through the compressor. Centrifugal compressors more nearly follow polytropic operation and are widely used to handle large volumes of gas at pressure ranges of 0.5 to several hundred psi. There are two ways to carry out the thermodynamic calculations for compression, namely by assuming:

1. An adiabatic (isentropic) reversible path: A process during which there is no heat added or removed from the system. The entropy is constant.

That is

$$pV^k = Constant$$
 (6-1)

2. A polytropic reversible path: A process in which changes in gas characteristics during compression are reviewed.

That is

$$pV^n = Constant$$
 (6-2)

The constant temperature process is a case when n=1, which is equivalent to isothermal compression, the constant pressure process n = 0; and the constant volume process $n = \infty$. Generally, it is impractical to build sufficient heat transfer equipment into the design of most compressors to convey the bulk of the heat of compression. Therefore most machines tend to operate along a polytropic path that approaches the adiabatic. Most compressor calculations are based on the adiabatic curve [3].

Relationships may be developed between the temperature, pressure, and volume for a polytropic process between state 1 and state 2. The ideal gas law states that:

$$pV = RT \tag{6-3}$$

That is

$$p_1 V_1^n = p_2 V_2^n \tag{6-4}$$

$$p_1 = \frac{RT_1}{V_1}, \quad p_2 = \frac{RT_2}{V_2}$$
 (6-5)

Rearranging p_1 and p_2 in Equation 6-5

$$\frac{\mathbf{p}_1}{\mathbf{p}_2} = \left(\frac{\mathbf{T}_1}{\mathbf{T}_2}\right) \left(\frac{\mathbf{V}_2}{\mathbf{V}_1}\right) \tag{6-6}$$

For a polytropic process between states 1 and 2,

$$\frac{\mathbf{p}_1}{\mathbf{p}_2} = \left(\frac{\mathbf{V}_2}{\mathbf{V}_1}\right)^n \tag{6-7}$$

Substituting Equation 6-6 into Equation 6-7

$$\left(\frac{\mathbf{T}_{1}}{\mathbf{T}_{2}}\right)\left(\frac{\mathbf{V}_{2}}{\mathbf{V}_{1}}\right) = \left(\frac{\mathbf{V}_{2}}{\mathbf{V}_{1}}\right)^{n}$$
(6-8)

Therefore

$$\frac{\mathbf{T}_1}{\mathbf{T}_2} = \left(\frac{\mathbf{V}_2}{\mathbf{V}_1}\right)^{n-1} \tag{6-9}$$

or

$$\frac{V_2}{V_1} = \left(\frac{T_2}{T_1}\right)^{\frac{1}{n-1}}$$
(6-10)

Substituting Equation 6-10 into Equation 6-7

$$\frac{p_1}{p_2} = \left(\frac{T_1}{T_2}\right)^{\frac{n}{n-1}}$$
(6-11)

The compression work can be calculated from the pressure-volume relationship as follows:

$$W = \int_{1}^{2} p dV$$
 (6-12)

and

 $pV^n = C$ for a polytropic process

$$W = \int_{1}^{2} C \frac{dV}{V^{n}}$$
(6-13)

Integrating Equation 6-13

$$W = C \int_{1}^{2} V^{-n} dv = \left[\frac{C V_{2}^{1-n} - C V_{1}^{1-n}}{1-n} \right]$$
(6-14)

$$\mathbf{C} = \mathbf{p}_2 \mathbf{V}_2^n = \mathbf{p}_1 \mathbf{V}_1^n$$

Therefore Equation 6-14 can be expressed as

$$\mathbf{W} = \begin{bmatrix} \underline{\mathbf{p}_2 \mathbf{V}_2 - \mathbf{p}_1 \mathbf{V}_1}\\ 1 - \mathbf{n} \end{bmatrix}$$
(6-15)

Using the ideal gas laws,

$$p_2V_2 = mRT_2$$

and

$$\mathbf{p}_1 \mathbf{V}_1 = \mathbf{m} \mathbf{R} \mathbf{T}_1$$

Equation 6-15 is given by

$$W = \frac{mR(T_2 - T_1)}{1 - n}$$
(6-16)

For the polytropic compression, work done is defined by

$$W = \frac{nR(T_2 - T_1)}{1 - n}$$
(6-17)

The polytropic compression work can be expressed as:

$$W = \frac{n}{n-1} [p_1 V_1 - p_2 V_2]$$
(6-18)

Since the process from 1 to 2 is polytropic, then

$$p_1 V_1^n = p_2 V_2^n$$
 (6-19)

and

$$\frac{V_2}{V_1} = \left(\frac{p_1}{p_2}\right)^{\frac{1}{n}}$$
(6-20)

From Equation 6-18

$$W = \frac{n}{n-1} p_1 V_1 \left[1 - \frac{p_2 V_2}{p_1 V_1} \right]$$
(6-21)

and

$$\frac{\mathbf{p}_{2}\mathbf{V}_{2}}{\mathbf{p}_{1}\mathbf{V}_{1}} = \left(\frac{\mathbf{p}_{2}}{\mathbf{p}_{1}}\right)^{\frac{n-1}{n}}$$
(6-22)

Substituting Equation 6-22 into Equation 6-21

$$W = \frac{n}{n-1} p_{i} V_{i} \left[1 - \left(\frac{p_{2}}{p_{1}} \right)^{\frac{n-1}{n}} \right]$$
(6-23)

where $R_c = p_2/p_1 = compression ratio$

Equation 6-23 is given by

$$W = \frac{n}{n-1} p_{I} V_{I} \left[1 - R_{c}^{\frac{n-1}{n}} \right]$$
(6-24)

Polytropic Compressor

Real compression processes operate between adiabatic and isothermal compression. Actual compression processes are polytropic processes. This is because the gas being compressed is not at constant entropy as in the adiabatic process, or at constant temperature as in the isothermal processes. Generally, compressors have performance characteristics that are analogous to those of pumps. Their performance curves relate flow capacity to head. The head developed by a fluid between states 1 and 2 can be derived from the general thermodynamic equation.

$$H = \int_{p_1}^{p_2} \overline{V} dP \tag{6-25}$$

where H = head, ft of fluid

p = pressure, psi $\overline{V} = specific volume of the fluid, ft^3/lb$

For polytropic compression, the pressure-volume relationship is

$$pV^n = constant$$

Compressors 427

$$V = \frac{C_1}{p^n}$$
(6-26)

where V = mole volume, ft³/(lb mol)

For the polytropic head, H_p , V can be substituted in Equation 6-25. The polytropic head is defined by

$$H_{p} = \int_{p_{1}}^{p_{2}} \frac{C_{1}}{p^{\frac{1}{n}}} dP$$
 (6-27)

Integrating Equation 6-27, H_p becomes

$$H_{p} = C_{1} \left(\frac{n}{n-1} \right) \left[p_{2}^{\frac{n-1}{n}} - p_{1}^{\frac{n-1}{n}} \right]$$
(6-28)

$$= C_{1} \left(\frac{n}{n-1}\right) p_{1}^{\left(\frac{n-1}{n}\right)} \left[\left(\frac{p_{2}}{p_{1}}\right)^{\frac{n-1}{n}} - 1 \right]$$
(6-29)

where

$$p_1^{\frac{1}{n}}V_1 = p_2^{\frac{1}{n}}V_2 = C_1$$
 and $R_c = \frac{p_2}{p_1}$ (6-30)

Substituting these into Equation 6-29, to eliminate C_1 gives

$$H_{p} = \left(\frac{n}{n-1}\right) p_{1} V_{1} \left[R_{c}^{\frac{n-1}{n}} - 1\right]$$
(6-31)

Using the gas law relationship

$$p_{1}V_{1} = \frac{Z_{1}RT_{1}}{M_{w}}$$
(6-32)

where Z_1 = compressibility factor at suction T_1 = absolute temperature at suction, °R M_w = molecular weight, (lb/lb mol) R = gas constant

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Substituting Equation 6-32 into Equation 6-31, the polytropic head, H_{p} , becomes:

$$H_{p} = \frac{Z_{1}RT_{1}}{M_{w}} \left(\frac{n}{n-1}\right) \left[R_{c}^{\frac{n-1}{n}} - 1\right]$$
(6-33)

If the compressibility factor, Z_{2} , for the gas at discharge conditions is significantly different from that of the suction, then the average compressibility factor, Z_{avg} is used to calculate the polytropic head.

$$Z_{avg} = \frac{Z_1 + Z_2}{2}$$
(6-34)

The polytropic head is defined by

$$H_{p} = \frac{Z_{avg}RT_{l}}{M_{w}} \left(\frac{n}{n-1}\right) \left[R_{c}^{\frac{n-1}{n}} - 1\right]$$
(6-35)

This can also be expressed as:

$$H_{p} = \left(\frac{1545. Z_{avg}. T_{1}}{M_{w}}\right) \left(\frac{n}{n-1}\right) \left[R_{c}^{\frac{n-1}{n}} - 1\right]$$
(6-36)

The discharge temperature, °F, is given by

$$t_{2} = \frac{(H_{p})(M_{w})}{(Z_{avg}.R)} \left(\frac{n-1}{n}\right) + t_{1}$$
(6-37)

There is a limit on the temperature as in the olefin or butadiene plants to prevent polymerization. At temperatures greater than $450 - 500^{\circ}$ F, the approximate mechanical limit, problems of sealing and casing growth could occur. High temperature requires a special, high machine cost. Therefore, multistage compressors are designed within the temperature range of 250° F - 300° F. In addition, a single-stage polytropic compressor can handle 7,000 to 11,000 feet of liquid, depending on the gas properties and inlet temperature [4].

The gas horsepower is expressed as

$$Gh_{p} = \frac{W.H_{p}}{1.98 \times 10^{6}.E_{p}}$$
(6-38)

The polytropic efficiency is used to compare adiabatic with polytropic performance. This is defined by

$$\frac{n}{n-1} = \left(\frac{k}{k-1}\right) \cdot E_p \tag{6-39}$$

Figure 6-2 shows the relationship between the polytropic efficiency and adiabatic (isentropic) efficiency of a perfect gas.

Adiabatic Compressor

The performance of reciprocating (piston) compressors with large valve area, or where valve losses are evaluated, is considered as close to

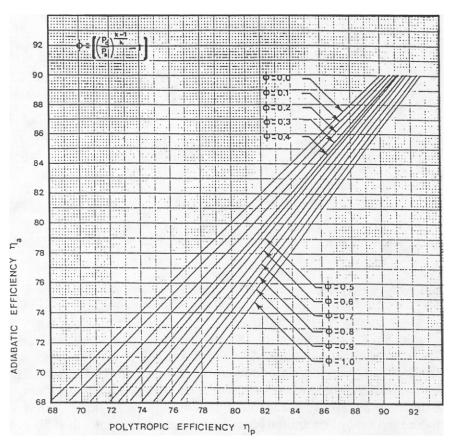


Figure 6-2. The relationship between the polytropic efficiency and the adiabatic efficiency for a perfect gas (Z = 1). (Source: A. K. Escoe, *Mechanical Design of Process Systems*, Volume 2, Gulf Publishing Company.)

adiabatic behavior as can be measured. The thermodynamic definition of an adiabatic process requires that no heat be added or removed from a system in which a change of state occurs. The adiabatic head produces the following equation, which is similar to the polytropic head of Equation 6-33. This is expressed as:

$$\mathbf{H}_{ad} = \left(\frac{\mathbf{Z}_{avg} \cdot \mathbf{R} \cdot \mathbf{T}}{\mathbf{M}_{w}}\right) \left(\frac{\mathbf{k}}{\mathbf{k}-1}\right) \left[\mathbf{R}_{c}^{\frac{\mathbf{k}-1}{\mathbf{k}}} - 1\right]$$
(6-40)

This can also be given by

$$H_{ad} = \left(\frac{1545. Z_{avg}. T_1}{M_w}\right) \left(\frac{k}{k-1}\right) \left[R_c^{\frac{k-1}{k}} - 1\right]$$
(6-41)

The discharge temperature is:

$$\mathbf{T}_{2} = \mathbf{T}_{1} \cdot \mathbf{R}_{c}^{\left(\frac{k-1}{k}\right)}$$
(6-42)

The adiabatic gas horsepower is:

$$Gh_{p} = \frac{W.H_{ad}}{1.98 \times 10^{6}.E_{ad}}$$
(6-43)

The adiabatic efficiency, E_{ad}, is:

$$E_{ad} = \frac{R_{c}^{\left(\frac{k-1}{k}\right)} - 1}{R_{c}^{\left(\frac{n-1}{n}\right)} - 1}$$
(6-44)

where

$$k = \frac{M_{w}C_{p}}{M_{w}C_{p} - 1.986}$$
(6-45)

Generally, if the compression ratio for piston machines is less than 5 or 6, single-stage compression is used. If the total compression ratio is between 6 and 36, two-stage compression will be required. Three or more stages may be required for compression ratios greater than 36. For two-stage compression, calculate the square root of the total compression ratio. Then assume that the compression ratio per stage will be equal to the square root of the total compression [2].

Mechanical Losses

After the gas horsepower is calculated by either the polytropic or adiabatic compression method, horsepower losses due to friction in bearings, seal, and speed increasing gears should be added. Although, there is no accurate method in estimating mechanical losses from gas power requirements, Table 6-1 gives approximate mechanical losses as a percentage of the gas power requirement [5].

The mechanical losses can be calculated by

Mechanical losses =
$$(Gh_p)(\%$$
 mechanical losses) (6-46)

The brake horsepower is:

 $Bh_p = Gh_p + mechanical losses$ (6-47)

The volumetric flow rate at suction or inlet condition can be expressed as:

$$Q_{s} = (ICFM) = \left[\frac{(MMSCFD) \times 10^{6}}{60 \times 24}\right] \left[\frac{14.7}{P_{s}}\right] \left[\frac{T_{s}}{520}\right] \\ \times \left[\frac{Z_{s}}{1.0}\right] \quad \text{ft}^{3}/\text{min}$$
(6-48)

The actual mass flow rate is

$$G = Q_{s} \left[\frac{P_{s} \cdot M_{w}}{1545Z_{s}T_{s}} \right] \quad lb/min$$
(6-49)

Table 6-1Approximate Mechanical Losses as a Percentageof a Gas Power Requirement

Gas English (hp)	Power Requirement Metric (kW)	Mechanical Losses %
0–3,000	0–2,500	3
3,0006,000	2,500-5,000	2.5
6,000-10,000	5,000-7,500	2
10,000+	7,500+	1.5

The accuracy of the developed polytropic head and efficiency curve may be reduced due to volume ratio effects. These effects are the influences that the performance of one impeller has on those of the impellers downstream. Lapina [1] stated these effects are greater as the number of impellers within a compression section increases. He proposed some guidelines to assess the volume-ratio effects for differences in the inlet conditions. An equation to calculate the value of θ for both the old or reference inlet conditions and the proposed new inlet conditions is given by

$$\theta = \left[\frac{(26.1)(M_{w}^{+})}{K.Z.T}\right]^{0.5}$$
(6-50)

MULTICOMPONENT GAS STREAMS

The design of a gas compressor for a gas mixture involves estimating the thermodynamic properties. The procedure for calculating the properties of a gas mixture is to use the weighted molal average of the property. These thermodynamic properties are estimated as follows:

Molecular weight
$$M_{w,mixture} = \sum_{i=1}^{n} y_i M_i$$
 (6-51)

Reduced temperature
$$T_{r,mixture} = \sum_{i=1}^{n} y_i T_{r,i}$$
 (6-52)

Reduced pressure
$$P_{r,mixture} = \sum_{i=1}^{n} y_i P_{r,i}$$
 (6-53)

Molal heat capacity
$$MC_{p,mixture} = \sum_{i=1}^{n} y_i MC_{p,i}$$
 (6-54)

Ratio of molal heat capacities
$$k_{mixture} = \frac{MC_{p,mixture}}{MC_{v,mixture}}$$

$$= \frac{MC_{p,mixture}}{(MC_{p,mixture} - 1.986)}$$
(6-55)

....

Compressibility factor $Z_{mixture} = f(T_r, P_r)$ for the mixture (6-56)

where $y_i =$ mole fraction of component i.

Table 6-2 summarizes the differences between reciprocating and centrifugal compressors.

Surge Control of Compressors

All dynamic compressors have a limited range of capacity for a given selection of impellers at a fixed speed. Below the minimum value of 50 to 70% of the rated flow, the compressor will surge, that is, it will become unstable in operation. Excessive vibration and possibly sudden failure or shutdown may arise.

It is essential that all compressor systems be designed to avoid possible surge operation. This is done by incorporating some type of antisurge control system. This is to ensure that the flow into the compressor is sufficient to maintain stability at the required pressure drop across the compressor. The surge control system should be used only for surge control and must not be linked with other functions of the process control system; otherwise, this could result in the damage of the compressor.

Centrifugal Compressors	Reciprocating Compressors
1. Lower installed first cost where pressure and volume conditions are favorable.	1. Greater flexibility in capacity and pressure range.
2. Lower maintenance cost.	2. High compressor efficiency and lower power cost.
3. Greater continuity of service and dependability.	3. Capability of delivering higher pressure.
4. Less operating attention.	4. Capability of handling smaller volumes.
5. Greater volume capacity per unit of plot area.	5. Less sensitive to changes in gas composition and density.
6. Adaptability to high speed low maintenance cost drivers.	

 Table 6-2

 Comparison of Reciprocating and Centrifugal Compressors

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Another surge control may be a blow off valve. This is automatically controlled to open or blow off excess capacity to the atmosphere if the process flow requirement is too low. In some cases, suction control valves are used. For gases that cannot be designed to atmosphere, a bypass control is the common antisurge control. This action bypasses unwanted flow back to the suction source. However, because the gas has already been compressed, its temperature is increased, and must therefore be cooled before entering the compressor a second time. In this case, a bypass cooler may be installed. The cooler may be avoided where the suction source is large or far away and the heat is dissipated by mixing or radiation. Figure 6-2A shows generalized flow characteristics of a compressor. In this figure, as the system resistance decreases, the compressor-system operating point moves to the right with a corresponding increase in the volumetric gas flow. However, the gas flow through the compressor cannot increase without a limit. This limit is known as "stonewall," and is caused by the choking of the gas as it exits from the compressor. These two operating limits are shown in Figure 6-2A. The designer should ensure that the system compressor curve-intersection point is well away from the surge or stonewall for efficient operation.

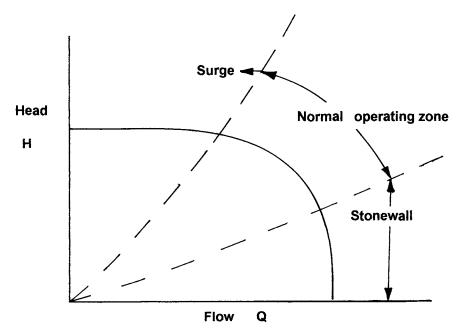


Figure 6-2A. Generalized flow characteristics of a compressor.

Treatment of Compressor Fluids

The discharge from any compressor is a dirty corrosive liquid. Removing the sludge from a compressor air system can effect the following:

- reduce installation costs incurred or drain traps, pipe and fittings, filters and regulators.
- reduce the maintenance of drain trap and the failure rate of pneumatic equipment, which is caused by dirt and moisture in the supply line.
- increase the life of pneumatic equipment.

Cases of high performance equipment require high quality of clean and dry compressed air. Compressed air contains water, oil, and dirty particles, which affects the performance of pneumatic equipment. Compressed air dryers are used to remove water vapor and to dry the air. Two main types of dryers are used in the CPI: refrigerant and desiccant dryers. Desiccant dryers use an adsorbent material, such as activated alumina, or molecular sieves to remove moisture from the compressed air. Also, desiccant dryers can be either heat regenerated or heatless. The main advantage of desiccant dryers is their ability to cool the air to a very low temperature. They are more expensive in terms of both capital and running costs. They are important when higher quality air is required. These dryers are widely used in offshore oil industry, where extreme ambient conditions are required. The design of desiccant dryers for removing water vapor from a natural gas is explained in Chapter Four.

Nomenclature

 $Bh_p = brake \text{ or shaft horsepower, bhp}$

- $\dot{C} = constant$
- C_p = specific heat at constant pressure, Btu/lb°F
- $\dot{C_v}$ = specific heat at constant volume, Btu/lb°F
- D = impeller diameter or rotor, ft
- $D_s =$ specific diameter, ft
- E_{ad} = adiabatic efficiency
- E_p = polytropic efficiency (try 0.75 for preliminary work)
- \dot{G} = gas flowrate, lb/min
- Gh_p = gas horsepower, actual compression horsepower, excluding mechanical losses, hp
 - $H = head, (ft. lb_f)/lb_m$

 H_{ad} = adiabatic head, (ft lb_f)/lb_m $H_n = polytropic head, (ft lb_f)/lb_m$ ICFM = volume flow referred to inlet condition, ft³/min $k = adiabatic (isentropic) exponent, C_p/C_v$ $MC_p = molar \text{ specific heat at constant pressure, Btu/(lb mol^{\circ}F)}$ $MC_v = molar \text{ specific heat at constant volume, Btu/(lb mol^{\circ}F)}$ M_{w} = molecular weight MMSCFD = million standard cubic feet per 24-hour day N = speed, rpm $N_s =$ specific speed, rpm n = polytropic exponentP = absolute pressure, psia P_1 = suction pressure, psia P_2 = discharge pressure, psia P_{c} = suction pressure, psia Q = volume flow, ft³/s Q_{s} = volume flow, ft³/min R = universal gas constant= 10.73(psia ft³/lb mole °R) = $1545[(lb ft^2) ft^3/lb mole ^R]$ or ft lb/lb mole ^R = $1.986(Btu/lb mole ^{\circ}R)$ $R_c = compression ratio, P_2/P_1$ T_1 = suction temperature, °R T_2 = discharge temperature, °R T_s = suction temperature, °R $t_1 = inlet temperature, ^{\circ}F$ t_2 = discharge temperature, °F W = workdone, Btu/lb-molew = gas flow, lb/h

 Z_{avg} = average compressibility factor for gas from suction to discharge conditions. A value of 1.0 will give conservative results.

CENTRIFUGAL PUMP HYDRAULICS

Introduction

Selecting the correct pump requires a knowledge of the system in which the pump must operate. In the chemical process industries (CPI), the engineer should start from the process flow sheets and schematics such as piping and instrument diagrams. The engineer must determine the optimum height and co-ordinate pump requirements, where pumps take suction from vessels or drums with the height above the pump. Where friction loss through piping forms a significant part of the total head, the designer should influence the selection of the allowable pressure drop (ΔP). In addition, volatile liquids, hot liquids, viscous liquids, and slurries will require a special approach and selection methods. If the pump is installed in a sump or a pit, essential factors require the correct sizing of the pit. These involve the flow requirements as the liquid flows toward the pump, and the positioning of the pump in the pit with sufficient baffles and spacing. Pump selection and troubleshooting are illustrated by Garay [6], McGuire [7], McNaughton [8], and have appeared in the literature [9,10,11].

This chapter considers the design of centrifugal pumps for given flow rates and physical properties.

Hydraulics of the System

The required factors in pump hydraulics are:

Density. The weight per unit volume of a substance, sometimes called specific weight. The density of water is 62.3 pounds per cubic foot.

Specific Gravity (**SpGr**). The ratio of the density or specific weight of a substance to that of some standard substance. For liquids, the standard is water usually at 60°F. In the petrochemical industry, API gravity use is 10° API. This corresponds to a specific gravity of 1.0.

SpGr(relative to water at 60° F) =
$$\frac{141.5}{131.5+° \text{ API}}$$
 (6-57)

In the chemical industry, degrees Baume' are commonly used. For liquids lighter than water,

$$SpGr = \frac{141.5}{130 + \text{ degrees Baume'}}$$
(6-58)

For liquids heavier than water,

$$SpGr = \frac{145}{145 - \text{ degrees Baume'}}$$
(6-59)

Pressure. The force exerted per unit area of a fluid, often designated as pounds per square inch (psi).

Barometric Pressure. The atmospheric pressure at the pump location. It changes with the weather conditions.

Gauge Pressure (psig). The gauge pressure is the pressure above atmospheric (or below in a vacuum).

Atmospheric Pressure (psi). The force exerted on a unit area by the weight of the atmosphere. The pressure at sea level due to the atmosphere is 14.7 psi.

Absolute Pressure (psia). The sum of atmospheric and gauge pressures. Absolute pressure in a perfect vacuum is 0, and at the atmosphere at sea level is 14.7 psi (0 psig).

1 atmosphere (bar) = 14.7 psi = 34 ft of water
conversion factor =
$$34/14.7 = 2.31$$

$$psi = \frac{head in feet \times SpGr}{2.31}$$
(6-60)

Vapor Pressure. The pressure at which a liquid at a specified temperature is in equilibrium with the atmosphere or with its vapor in a closed system. At pressures below the vapor pressure at a given temperature, the liquid will start to vaporize. This is due to a reduced pressure at the surface of the liquid.

Head. The height of a fluid column, measured in feet, to ascertain the gauge pressure at the bottom of the column.

Head, ft =
$$\frac{2.31(\text{pressure, psi})}{(\text{SpGr})}$$
 (6-61)

Friction Head. The pressure to overcome the resistance to the flow in the pipe and fittings. It is expressed as lbs/sq. in. or feet of liquid.

Static Head. Fluid elevation above or below the datum line. For horizontal centrifugal pumps, static head is calculated from the pump center line; with vertical pumps, it is taken from the eye of the first-stage impeller.

Velocity Head. The kinetic energy of a flowing fluid. It is defined by

$$k = \frac{V^2}{2g}$$
(6-62)

Suction Lift. The manometer reading in feet of liquid at the pump suction (corrected to a datum), minus the velocity head. This term expresses the suction head when it is below atmospheric pressure.

Static Suction Lift. The vertical distance from the center line of the pump down to the free level of the liquid source.

Net or Dynamic Suction Lift. Static suction lift, friction head, and velocity head.

Suction Head. Occurs when the source of supply is above the center line of the liquid.

Static Discharge Head. The vertical elevation from the center line of the pump to the point of free discharge.

Dynamic Discharge Head. The static discharge head plus dynamic suction lift or minus dynamic suction head.

Total Discharge Head. The pressure reading at the pump discharge in feet of liquid, corrected to a datum, plus the velocity head.

Total Head. The algebraic difference between the discharge and suction heads. A suction lift is added to the total discharge head; a positive suction head is subtracted from the discharge head.

Head Losses. The frictional head losses caused by fluid flow through pipes, fittings, valves, and nozzles.

NPSH. The net positive suction head is the most critical factor in a pumping system. A sufficient NPSH is essential, whether working with centrifugal, rotary, or reciprocating pumps. Marginal or inadequate NPSH will cause cavitation, which is the formation and rapid collapse of vapor bubbles in a fluid system. Collapsing bubbles place an extra load on pump parts and can remove a considerable amount of metal from impeller vanes. Cavitation often takes place before the symptoms become evident. Factors that indicate cavitation are increased noise, loss of discharge head, and reduced fluid flow.

Pump Hydraulics Calculations

Pump sizing computation requires the following information on the process flow diagram or a pump calculation sheet: service, size and type, fluid, pump temperature, density at pump temperature, design capacity in gpm at pump temperature, suction head, discharge head, design differential, NPSH (net positive suction head) and power requirements.

Calculating the suction condition of a pump comprises the following:

- 1. vessel pressure (original pressure).
- 2. add the suction head or minus for a suction lift, S.Hd.
- 3. minus the line pressure drop, L loss.

That is

Suction pressure = orig. pres. + S.Hd. - L. loss (6-63)

The discharge condition of a pump considers the following:

- 1. delivery pressure
- 2. static head
- 3. control valve pressure drop
- 4. exchanger pressure drop
- 5. orifice pressure drop
- 6. furnace pressure drop
- 7. line pressure drop
- 8. other pressure drop

The NPSH = (orig. pres. - vapor pres.) + S.Hd - L. loss, ft (6-64)

The net head at the suction of the pump impeller must not exceed a certain value to prevent formation of vapor and cause cavitation of the metal. This minimum head is called the available $NPSH_a$, and must be greater than the required $NPSH_r$ for a stable pump operation.

Pump discharge pres. = del. pres. + Shd.
+
$$\Delta P$$
 Cont. v. + ΔP exch.
+ ΔP orif. + ΔP furn.
+ ΔP line + ΔP Other. (6-65)

Total pump ΔP = pump discharge pres. – pump suction pres. (6-66)

Bhp (brake horsepower) =
$$\frac{(\text{flow rate, gpm})(\text{pump }\Delta \text{P, psi})}{(1715)(\text{EFF})}$$
(6-67)

Bhp =
$$\frac{(\text{gpm})(\text{H})(\text{SpGr})}{(3960)(\text{EFF})}$$
 (6-68)

where H = total discharge head

Table 6-3 shows a pump hydraulic design calculation data sheet.

Centrifugal Pump Efficiency

The design engineer must use the expected pump efficiency provided in the pump performance curve to evaluate the required brake horsepower for a centrifugal pump. In the early stages of the design, it is customary to estimate a value for the efficiency. Final values depend on the specified pump and the operating conditions that are encountered. Branan [4] has developed an equation to calculate the centrifugal pump efficiency and pump horsepower. The equation was based on pump efficiency curves of the NGPSA Engineering Data Book. These efficiency

Fluid		Gas Oil	
Viscosity at Pumping temperature (P.T), cP	0.6	
Vapor Pressure at pumping tempera (P.T), psia	iture	0.01	
Specific gravity, (SpGr) at P.T.		1.04	
Density at P.T., lb/ft ³		64.87	
Operating flow at P.T., gpm		250.0	
Design Flow at P.T., gpm		250.0	
Suction		Discharge	
Vessel pressure, Orig. Pres., psia	27.7	Delivery pressure, psia	a. 28.2
Static head S.Hd., psi	5.4	Static head S.Hd., psi	14.86
– Line loss ΔPf, psi.	0.23	ΔP Control valve, psi	10.0
Suction pres. = Orig. pres. + S.Hd. - L. line	32.78	ΔP Exchanger, psi	5.0
NPSH = (orig. pres. – vapor pres.) · S.Hd. – L.loss, psia.	+ 32.77	ΔP Orifice, psi ΔP Furnace , psi	2.0
		Line loss ΔP psi	4.38
		Other ΔP , contigency,	psi

Table 6-3Pump Hydraulic Design Calculation Sheet

curves give good results with the vendor data (the range of developed heads, 50ft to 300ft, and the flow rates, 100 to 1000 gpm). The equation for calculating the pump efficiency is expressed as:

$$EFF = (80 - 0.2855H + 3.78 \times 10^{-4} HQ)$$

- 2.38 × 10⁻⁷ HQ² + 5.39 × 10⁻⁴ H²
- 6.39 × 10⁻⁷ H²Q + 4 × 10⁻¹⁰ H²Q²)(1/100) (6-69)

The actual brake horsepower required for pump operation Bhp is:

$$Bhp = \frac{HBhp}{EFF}$$
(6-70)

where EFF = pump efficiency (decimal fraction)

H = developed pump head, ft

- Q = flow rate through pump, gpm
- HBhp = hydraulic brake horsepower, hp

Bhp = actual brake horsepower required for pump operation, hp

Using Equation 6-69, Figure 6-3 shows typical characteristic curves for a 6-inch centrifugal pump operating at 1750 rpm. Equation 6-69 gives

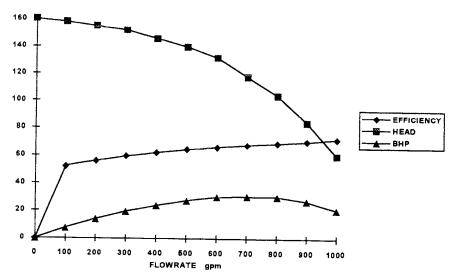


Figure 6-3. Typical characteristic curves for a 6-inch centrifugal pump operating at 1750 rpm.

results within 7% of the actual pump curves for the range of applicability (H = 50 to 300 ft, Q = 100 to 1000 gpm). For flows in the range of 25 to 99 gpm, an approximate efficiency can be obtained by solving Equation 6-69 for 100 gpm and then subtracting 0.35% gpm times the difference between 100 gpm and the low flow rate in gpm. For low flow rates, near 25–30 gpm, this will give results within about 15% for the middle of the head range (pump ΔP) and 25% at the extremes. The horsepower at the 25–30 gpm level is normally below 10 [4].

AFFINITY LAWS

The affinity laws express the mathematical relationship between the variables involved in a pump's performance. By applying the principles of dimensional analysis on the physical properties affecting pump operation, the relationship is:

$$f\left\{\frac{Q}{(g_{c}H)^{0.5}D^{2}}, \frac{NQ^{0.5}}{(g_{c}H)^{0.75}}, \frac{\rho QD}{\mu D^{2}}\right\} = 0$$
(6-71)

where the dimensionless expressions are:

$$\pi_1 = \frac{Q}{(g_c H^{0.5}) D^2}$$
(6-72)

$$\pi_2 = \frac{NQ^{0.5}}{(g_c H)^{0.75}}$$
(6-73)

$$\pi_3 = \frac{\rho Q D}{\mu D^2} \tag{6-74}$$

For true dynamic similarity, π_1 , π_2 , and π_3 must be constant for similar pumps or for the same pump at different speeds. π_3 , which is the Reynolds number, cannot be maintained constant with changes in pump speed or size if the same liquid is pumped. Therefore, for practical application, the Reynolds number can be neglected when considering the conditions for dynamic similarity.

The conditions for dynamic similarity are constant values of π_1 and π_2 . Equation 6-71 can be altered by appropriate mathematical operation to yield

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$$f\left\{\frac{Q}{(g_{c}H)^{0.5}D^{2}}, \frac{ND}{(g_{c}H)^{0.5}}\right\} = 0$$
(6-75)

where

$$\pi_2' = \frac{\text{ND}}{(g_c H)^{0.5}} \tag{6-76}$$

Therefore, for conditions of similarity of flow π_1 , π_2 , and π'_2 are constant. Stepanoff [12] has shown that the velocity diagrams at the impeller are geometrically similar when the pumps are operating at the same efficiency for similar pumps or for the same pump at different speeds.

Applications of Affinity Laws

For the same pumps or geometrically similar pumps, $Q/(g_cH)^{0.5}D^2$, $NQ^{0.5}/(g_cH)^{0.75}$ and $ND/(g_cH)^{0.5}$ have unique values corresponding to each efficiency. In addition, these values remain constant despite the impeller diameter or pump speed. For the same pump operating at different speeds the impeller diameter, D, is constant.

That is

D = constant

Therefore at any given efficiency

Capacity:

$$\frac{Q}{(g_c H)^{0.5} D^2}$$
 and $\frac{ND}{(g_c H)^{0.5}}$ are constant

or

$$\frac{Q}{(g_{c}H)^{0.5}D^{2}} / \frac{ND}{(g_{c}H)^{0.5}} = Constant$$
(6-77)

Therefore

$$\frac{Q}{ND^3} = Constant = K$$
(6-78)

But D is constant Therefore

Q = K N (6-79)

where K' and K are constants Therefore

$$\frac{Q_2}{Q_1} = \frac{N_2}{N_1}$$
(6-80)

Head:

$$\frac{\text{ND}}{(g_c H)^{0.5}} = \text{Constant} = J$$
(6-81)

or

$$\frac{\mathrm{N}^2\mathrm{D}^2}{\mathrm{H}} = \mathrm{J}' \tag{6-82}$$

where J and J' are constants.

Therefore at a constant diameter

$$\frac{\mathrm{H}_2}{\mathrm{H}_1} = \left(\frac{\mathrm{N}_2}{\mathrm{N}_1}\right)^2 \tag{6-83}$$

Break horsepower (Bhp)

Bhp α QH α N³ at constant D

Therefore

$$\frac{Bhp_2}{Bhp_1} = \left(\frac{N_2}{N_1}\right)^3 \tag{6-84}$$

These laws apply only for efficiencies of the same value.

For geometrically similar pumps with different impeller diameters but same speeds:

From Equation 6-78

$$Q = KND^3$$
(6-85)

At constant speed

 $Q = K^{"}D^{3}$ where K and K" are constants.

$$\frac{\mathbf{Q}_2}{\mathbf{Q}_1} = \left(\frac{\mathbf{D}_2}{\mathbf{D}_1}\right)^3 \tag{6-86}$$

Head:

At constant speed from Equation 6-82

$$\frac{\mathrm{H}_2}{\mathrm{H}_1} = \left(\frac{\mathrm{D}_2}{\mathrm{D}_1}\right)^2 \tag{6-87}$$

Brake horsepower (Bhp)

$$\frac{Bhp_2}{Bhp_1} = \left(\frac{D_2}{D_1}\right)^5$$
(6-88)

Specific Speed

One dimensionless expression in the affinity laws is:

 $\frac{NQ^{0.5}}{(g_cH)^{0.75}} = \text{constant at a given efficiency}$

It is often used by designers in sizing and selecting the type of pump or compressor. The performance of centrifugal pumps is related to a parameter called specific speed. It is a correlation of capacity, head, and speed at optimum efficiency that classifies pump impellers with respect to their geometric similarity. Specific speed is a number expressed algebraically as:

$$N_{s} = \frac{NQ^{0.5}}{H^{0.75}}$$
(6-89)

where N_s = specific speed of the pump

N = pump speed, rpm

Q = flow at optimum efficiency, gpm

H = head per impeller, ft

The specific speed of an impeller is the rotation in rpm. This is a geometrically similar impeller that would run if it were of such size as to discharge 1 gpm against a total head of 1 ft. Specific speed is used to determine the NPSH required by a given impeller. Those that produced high total dynamic head (TDH) have low specific speeds, and conversely, low TDH impellers have high specific speeds. At a particular head and capacity, a pump of low specific speed will operate safely at lower NPSH than one with a high specific speed. Designers should ensure that the capacity or TDH is increased when recommending speeding up an existing pump. This will enable that the new NPSH does not exceed what is available.

$$D_{s} = \frac{DH^{0.25}}{Q^{0.5}}$$
(6-90)

Figure 6-4 shows the dimensionless parameters as presented by Balje. The figure is the graphical combination of Equations 6-89 and 6-90.

Nomenclature

Bhp = brake or shaft horsepower, bhp $C = constant$
C = constant
D = impeller diameter or rotor, ft
del. pres. = pump delivery pressure, psia
$D_s = specific diameter, ft$
EFF = pump efficiency
gpm = fluid flow rate through pump
H = head, (ft lb)/lb
HBhp = hydraulic brake horsepower required by pump (pump
efficiency = 1.0 for HBHP calculation)
L. loss = pressure drop due to piping, psi or ft
orig. pres. = vessel pressure on pump suction, psia
pump dish. pres. = pump discharge pressure, psia
Q = Volumetric flowrate, gpm
vap. pres. = vapor pressure of pumped fluid at pumping tempera-
ture, psia
suc. pres. = pump suction pressure, psia
S.Hd. = static head of liquid on pump suction or discharge, ft
total pump ΔP = pressure rise across pump, psi or ft

PROBLEMS AND SOLUTIONS

Problem 6.1

A centrifugal compressor is to be specified for a gas plant. The unit is to compress 18,000 lb_m/hr of gas mixture at 50 psia and 150°F to 200 psia. The gas mixture consists of 30% hydrogen (H₂), 45% methane (CH₄),

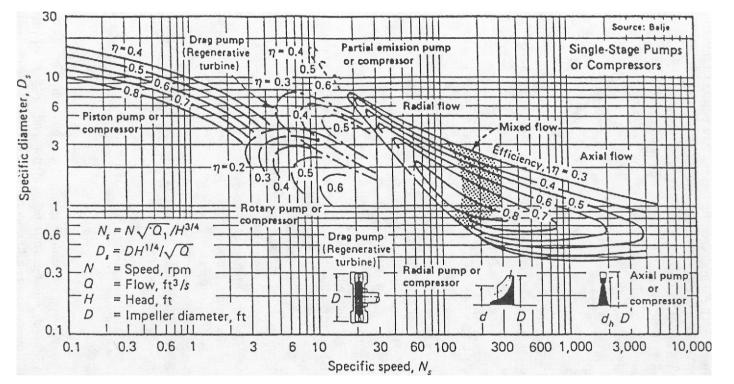


Figure 6-4. Specific speed and specific diameter enable the initial selection of a definite type of single-stage compressor. (O. E. Balje, "A Study on Design Criteria and Matching of Turbomachines: Part B", Transaction of the American Society of Mechanical Engineers, *Journal of Engineering Power*, Vol. 84, January 1962.)

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15% ethane (C₂H₆), 7% propane (C₃H₈), and 3% n-butane (n – C₄H₁₀). Assuming a polytropic efficiency of 75%, and the average compressibility factor, $Z_{avg} = 0.95$, calculate the polytropic head, total break horsepower, and the discharge temperature. Table 6-4 gives the properties of the gas mixture.

The mixture specific heat ratio, k, is defined by

$$k = \frac{MC_{p}}{MC_{v}} = \frac{MC_{p}}{MC_{p} - 1.986}$$
$$k = \frac{10.317}{8.331}$$
$$k = 1.238$$

The average molecular weight, $M_w = 17.162$

Solution

The computer program PROG61 calculates the polytropic and adiabatic heads of a compressor. Table 6-5 shows the input data and results of the gas mixture. The gas discharge temperature is 310.4° F and the compressor polytropic head is 76874 ft with a total break horsepower \cong 960 hp.

		Prope	rties of ti	ne Gas M	lixture		
Gas	Mole fraction y	Molecular weight M _w	уМ _w	P _c psia	T _c (°R)	$ \begin{array}{c} MC_{p} \\ \left(\frac{Btu}{Ib_{m} \cdot mol.^{\circ}R} \right) \\ @ 150^{\circ}F \end{array} $	уМС _р @ 150°F
H_2	0.30	2.01	0.603	188.1	60.2	6.94	2.082
CH_4	0.45	16.04	7.218	666.0	343	8.95	4.028
C_2H_6	0.15	30.07	4.511	707	550	13.78	2.067
C_3H_8	0.07	44.09	3.086	616	660	19.52	1.366
nC_4H_{10}	0.03	58.12	1.744	551	765	25.81	0.774
		M _w =	17.162			MC _p =	10.317

Table 6-4 Properties of the Gas Mixture

Table 6-5
Input Data and Computer Output of Polytropic
Compressor Calculations

DATA61.C	DAT		
POLYTROP	PIC		
18000.0	50.0	200.0	
		17.162	
0.95	0.75		
		POLYTROPIC COMPRESSOR CAL	CULATION
******	******	******	*****
TYPE	C OF SERV	ICE:	POLYTROPIC
		., lb/hr.:	18000.000
SUCI	TION PRES	SURE, psia:	50.000
DISC	CHARGE PR	RESSURE, psia:	200.000
COMP	RESSION	RATIO:	4.000
INLE	T TEMPER	ATURE, OF:	80.000
RATI	O OF SPE	CIFIC HEAT CAPACITIES, k:	1.238
MOLE	CULAR WE	IGHT OF GAS, lb/lb-mole:	17.162
AVEF	RAGE COMP	PRESSIBILITY FACTOR, Z:	.9500
POLY	TROPIC E	CFFICIENCY:	.75
POLY	TROPIC H	IEAD, ft:	76873.770
GAS	DISCHARC	E TEMPERATURE, oF:	310.404
GAS	HORSEPOW	IER, hp:	931.803
MECH	HANICAL L	OSSES, hp:	27,954
TOT	L BRAKE	HORSEPOWER, hp:	959.757
WORK	CONE, 1	b-mole/hr.:	735.634

Problem 6.2

Size a centrifugal pump with a 4-inch suction nozzle, and a 3-inch discharge nozzle that handles a gas oil at normal flow rate of 250 gpm through piping and component's system as shown in Figure 6-5, at the following conditions, temperature $t = 555^{\circ}F$, specific gravity = 1.04, and viscosity of the gas oil $\mu = 0.6$ cP. Table 6-3 gives the details on a pump calculation sheet. Other piping details are shown in Table 6-6.

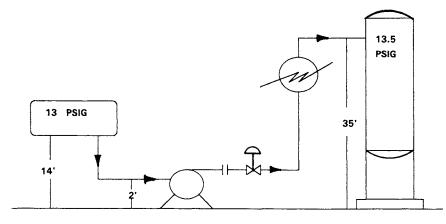


Figure 6-5. Gas oil transfer pump system.

Schedule 40	Suction Line	Discharge Line
Normal size, inch	6	4
Inside diameter, inch	6.065	4.026
Actual pipe length,ft	39	156
Fittings		
Long radius 90° elbows	5	20
Reducer, inch	4.026	3.068
Gate valve	1	4
Entrance	1	
exit		1

Table 6-6 Piping Details and Fittings

Solution

The line pressure drops for the suction and the discharge can be calculated by the method described in Chapter Three.

The total suction pressure drop is:

Total suction $\Delta P_{suct} = 5.40 + 27.7 - 0.23$

= 32.87 psi

Total discharge $\Delta P_{disch} = 28.2 + 14.86 + 10.0 + 5.0 + 2.0 + 4.38$

= 64.44 psi

Differential pressure $\Delta P = \Delta P_{disch} - \Delta P_{suct}$

The brake horsepower Bhp = $\frac{(Q, gpm)(\Delta P, psi)}{(1715)(EFF)}$

$$= \frac{(250)(31.57)}{(1715)(0.7)}$$
$$= 6.57 \text{ hp}$$
$$\cong 7.0 \text{ hp}$$

The computer program PROG62 sizes the centrifugal pump for the given flow rate and fluid characteristics. The program calculates the hydraulic brake horsepower required by the pump and the actual brake horsepower. In addition, the program computes the available net positive suction head (NPSH) and the pump efficiency. Table 6-7 shows the input data and results of the pump hydraulic design calculation. The available NPSH is 64 ft, and the actual brake horsepower required for the pump operation is 7.0hp, with a pump efficiency of 68%.

 Table 6-7

 Centrigugal Pump Sizing: Program Inputs and Outputs

DATA62.DAT	
GAS-OIL	
555.0 0.6 1.04	
4.0 250.0	
27.7 0.22 12.0	
28.2 33.0 10.0	
5.0 2.0 0.0	
4.38 0.0 0.0	
CENTRIFUGAL PUMP SIZING	
* * * * * * * * * * * * * * * * * * * *	
FLUID NAME:	GAS-OIL
FLUID OPERATING TEMPERATURE, oF:	555.000
FLUID VISCOSITY, CP:	.6000
FLUID SPECIFIC GRAVITY:	1.040
FLUID VAPOR PRESSURE, psia:	4.000
VOLUMETRIC FLOWRATE:, gpm:	250.000
PRESSURE AT EQUIPMENT, psia:	27.700
SUCTION LINE LOSSES, psi:	.220
OTHER LOSSES (e.g. STRAINER/FILTER), psi:	.000
SUCTION STATIC HEAD, ft.:	12.000
SUCTION STATIC HEAD, psi:	5.403
TOTAL PUMP SUCTION PRESSURE, psig:	18.183
TOTAL PUMP SUCTION PRESSURE, psia:	32.883
AVAILABLE NET POSITIVE SUCTION HEAD, ft.:	64.153
DELIVERY PRESSURE, psia.:	28.200
DISCHARGE STATIC HEAD, ft.:	33.000
DISCHARGE STATIC HEAD, psi:	14.857
CONTROL VALVE PRESSURE DROP, psi.:	10.000
EXCHANGER PRESSURE DROP, psi.:	5.000
ORIFICE PRESSURE DROP, psi.:	2.000
FURNACE PRESSURE DROP, psi.:	.000
DISCHARGE LINE LOSSES, psi.:	4.380
OTHER LOSSES, psi.:	.000
TOTAL PUMP DISCHARGE PRESSURE, psig:	49.737
TOTAL PUMP DISCHARGE PRESSURE, psia:	64.437
TOTAL PUMP PRESSURE DROP, psi.:	31.555
TOTAL PUMP PRESSURE DROP, ft.:	70.088
HYDRAULIC BRAKE HORSEPOWER REQUIRED BY THE PUMP, bhp:	4.600
ACTUAL BRAKE HORSEPOWER REQUIRED FOR PUMP OPERATION bhp:	
PUMP EFFICIENCY, (%):	67.557

Problem 6.3

A 6-inch pump delivers the following heads when rotating at 1750 rpm and at varying flow rates as shown in Table 6-8. For a geometrically similar 8-inch pump running at 1450 rpm, calculate:

- 1. the heads developed by the 8-inch pump.
- 2. the brake horsepower for the flow rates.
- 3. the pump efficiencies for the 6- and 8-inch pumps.
- 4. develop the characteristic curves for the 6-inch and the 8-inch pumps.

Solution

Using the affinity law (Equation 6-76), the corresponding values for the head at varying flow rates are shown in Table 6-8. The computer program PROG63 calculates both the brake horsepower and pump efficiencies of the 6- and 8-inch pumps. Tables 6-9 and 6-10 show the input data and the computer results of these pumps. The characteristic curves are shown in Figure 6-6.

6 inch @ Q (gpm)	1750 rpm H (ft)	8 inch @ Q (gpm)	1450 rpm H (ft)
0	160	0	195.3
100	158	100	192.8
200	155	200	189.2
300	152	300	185.5
400	146	400	178.2
500	140	500	170.9
600	132	600	161.1
700	118	700	144.0
800	104	800	126.9
900	84	900	102.5
1000	60	1000	73.2

Table 6-8 Total Heads and Flow Rates for the 6-inch and 8-inch Centrifugal Pumps

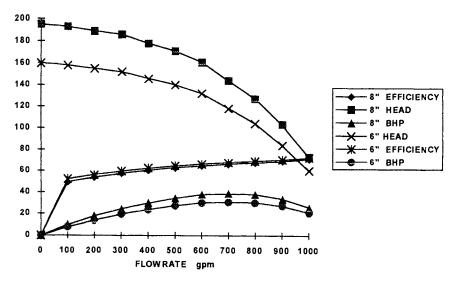


Figure 6-6. Characteristic curves of the 6-inch and the 8-inch centrifugal pumps.

Table 6-9 Efficiency and Brake Horsepower Calculations for the 6-inch Centrifugal Pump Program Inputs and Outputs

DATA63.DAT			
11 0.0 160.0			
100.0 158.0			
200.0 155.0			
300.0 152.0			
400.0 146.0			
500.0 140.0			
600.0 132.0			
700.0 118.0			
800.0 104.0			
900.0 84.0			
1000.0 60.0			
	EFFICIENCY AN	D BRAKE HORSEPOWER	CALCULATIONS
*****			*****
FLOWRATE gpm	HEAD It.	EFFICIENCY *	BRAKE HORSEPOWER Bhp
	160.0	~	.0
.0 100.0	158.0	.0 52.4	7.6
200.0	155.0	56.3	13.9
300.0	152.0	59.4	19.4
400.0	146.0	62.2	23.7
500.0	140.0	64.4	27.4
600.0	132.0	66.2	30.2
700.0	118.0	67.8	30.8
800.0	104.0	69.0	30.5
900.0	84.0	70.4	
1000.0	04.0	/0.4	27.1
1000.0	60.0	72.3	27.1 20.9

DATA63.DAT			
11			
0.0 195.3			
100.0 192.8 200.0 189.2			
300.0 185.5			
400.0 178.2			
500.0 170.9			
600.0 161.1			
700.0 144.0			
800.0 126.9			
900.0 102.5 1000.0 73.2			
****		ID BRAKE HORSEPOWER	CALCULATIONS
**************************************	**************************************		
FLOWRATE gpm	**************************************	EFFICIENCY %	BRAKE HORSEPOWER Bhp
FLOWRATE gpm	+*************************************	EFFICIENCY %	BRAKE HORSEPOWER Bhp ************************************
FLOWRATE gpm ************************************	HEAD ft. ++++++++++++++++++++++++++++++++++++	EFFICIENCY % .0 49.6 53.8	BRAKE HORSEPOWER Bhp .0 9.8 17.8
FLOWRATE gpm ************************************	HEAD ft. 195.3 192.8 189.2 185.5	EFFICIENCY % .0 49.6 53.8 57.3	BRAKE HORSEPOWER Bhp ************************************
FLOWRATE gpm ************************************	HEAD ft. ++++++++++++++++++++++++++++++++++++	EFFICIENCY % ************************************	BRAKE HORSEPOWER Bhp ************************************
FLOWRATE gpm ************************************	HEAD ft. ************************************	EFFICIENCY % ************************************	BRAKE HORSEPOWER Bhp ************************************
FLOWRATE gpm ************************************	HEAD ft. 195.3 192.8 189.2 185.5 178.2 170.9 161.1	EFFICIENCY % ************************************	BRAKE HORSEPOWER Bhp .0 9.8 17.8 24.5 29.8 34.4 37.8
FLOWRATE gpm ************************************	HEAD ft. ++++++++++++++++++++++++++++++++++++	EFFICIENCY % ************************************	BRAKE HORSEPOWER Bhp ************************************
FLOWRATE gpm ************************************	HEAD ft. 195.3 192.8 189.2 185.5 178.2 170.9 161.1	EFFICIENCY % ************************************	BRAKE HORSEPOWER Bhp .0 9.8 17.8 24.5 29.8 34.4 37.8

Table 6-10 Efficiency and Brake Horsepower Calculations for the 8 inch Centrifugal Pump Program Inputs and Outputs

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PROGRAM PROG61

000000000	THIS PROGRAM CALCULATES EITHER POLYTROPIC OR ADIABATIC HEADS, THE ASSOCIATED GAS HORSEPOWER AND GAS DISCHARGE TEMPERATURE, AS LONG AS THE POLYTROPIC OR ADIABATIC EFFICIENCY IS USED WITH THE COMPANION HEAD CALCULATION. THE PROGRAM ALSO CALCULATES ADIABATIC EFFICIENCY FROM POLYTROPIC EXPONENT "K" FROM THE MOLAL HEAT CAPACITY (MCp).
с с	***************************************
0 0 0 0 0 0 0 0	Hpoly(PH) = POLYTROPIC HEAD, ft ZAV = AVERAGE COMPRESSIBILITY FACTOR FOR GAS FROM SUCTION TO DISCHARGE CONDITIONS. A VALUE OF 1.0 WILL GIVE CONSERVATIVE RESULTS. R = GAS CONSTANT= 1544. TS = SUCTION TEMPERATURE, OF MW = MOLECULAR WEIGTH OF GAS
с с с с	N = POLYTROPIC EXPONENT RC = COMPRESSION RATIO (DISCHARGE PRESSURE/SUCTION PRESSURE) P2/P1 T11 = INLET TEMPERATURE, oF
	<pre>T12 = DISCHARGE TEMPERATURE, oF K = ADIABATIC EXPONENT, Cp/CV Epoly (PE) = POLYTROPIC EFFICIENCY (TRY 0.75 FOR PRELIMINARY WORK) HP = GAS HORSEPOWER, bhp. W = GAS FLOW, 1b/h Had (AH) = ADIABATIC HEAD, ft T21 = SUCTION TEMPERATURE, oF Ead (AE) = ADIABATIC EFFICIENCY MCp (MCP) = MOLAL SPECIFIC HEAT OF GAS AT CONSTANT PRESSURE, Btu/(1b mol)(oF). P1 = SUCTION PRESSURE, psia. P2 = DISCHARGE PRESSURE, psia. P2 = DISCHARGE PRESSURE, psia. CHOICE = CHOOSE WHETHER THE PROCESS IS ADIABATIC OR POLYTROPIC ************************************</pre>
	COMMON/PRÓG1/W,P1,P2,T11,T12,T21,T22,K,MW,N,ZAV,AE,AECAL,RC COMMON/PROG2/POLYH,ADIAH,HPPOLY,HPADIA,PE,PECAL,BHPP,BHPA COMMON/PROG3/WORK,PROD
	OPEN (UNIT=3,FILE='DATA61.DAT',STATUS='OLD',ERR=18) OPEN (UNIT =1, FILE ='PRN')
c c c	DETERMINE WHETHER THE COMPRESSOR CALCULATION IS POLYTROPIC OR ADIABATIC.
	READ (3, 15, ERR=19) CHOICE
15	FORMAT (A)
	IF (CHOICE .EQ. 'POLYTROPIC' .OR. CHOICE .EQ. 'polytropic') THEN

```
READ (3, *, ERR=19) W, P1, P2
READ (3, *, ERR=19) T11, K, MW
READ (3, *, ERR=19) ZAV, PE
           GO TO 40
       ELSEIF (CHOICE .EQ. 'ADIABATIC' .OR. CHOICE .EQ. 'adiabatic') THEN
           READ (3, *, ERR=19) W, P1, P2
READ (3, *, ERR=19) T11, K, MW
READ (3, *, ERR=19) ZAV, AE
           GO TO 50
           ENDIF
           WRITE (*, 10)
18
           FORMAT (3X, 'FILE DOES NOT EXIST')
10
           GO TO 999
          WRITE (*, 20)
FORMAT (3X, 'ERROR MESSAGE IN THE DATA VALUE')
19
20
           GO TO 999
40
           WRITE (1, 30)
           FORMAT (///,26X,'POLYTROPIC COMPRESSOR CALCULATION', /, 78(1H*))
30
           CALL POLYTR
          WRITE (1, 60) CHOICE, W, P1, P2, RC
FORMAT (5X,'TYPE OF SERVICE:', T60, A,/,
5X,'GAS FLOWRATE, lb/hr.:',T60, F12.3,/,
5X,'SUCTION PRESSURE, psia:', T60, F12.3,/,
5X,'DISCHARGE PRESSURE, psia:', T60, F12.3,/,
5X,'COMPRESSION RATIO:', T60, F12.3)
60
      *
      *
      ×
      ×
          70
      *
                        5X,'MOLECULAR WEIGHT OF GAS, lb/lb-mole:',T60,F12.3)
      ٠
           WRITE (1, 80) ZAV, PE
          FORMAT (5X,'AVERAGE COMPRESSIBILITY FACTOR, Z:',T60, F8.4,/,
5X,'POLYTROPIC EFFICIENCY:',T60,F8.2)
80
           WRITE (1, 90) POLYH, T12, HPPOLY
           FORMAT (5X,'POLYTROPIC HEAD, ft:',T60,F12.3,/,
5X,'GAS DISCHARGE TEMPERATURE, oF:',T60, F12.3, /,
5X,'GAS HORSEPOWER, hp:', T60, F12.3)
90
      *
          WRITE (1, 100) PROD, BHPP, WORK
FORMAT (5X,'MECHANICAL LOSSES, hp:', T60, F12.3,/,
5X,'TOTAL BRAKE HORSEPOWER, hp:',T60, F12.3,/,
5X,'WORK DONE, lb-mole/hr.:', T60, F12.3)
100
      +
           WRITE (1, 105)
```

105	FORMAT (78(1H-))
с	FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR(12)
	GO TO 999
50 110	WRITE (1, 110) FORMAT (///,26X,'ADIABATIC COMPRESSOR CALCULATION', /, 78(1H*))
	CALL ADIAB
120 * * *	<pre>WRITE (1, 120) CHOICE, W, P1, P2, RC FORMAT (5X,'TYPE OF SERVICE:', T60, A,/, 5X,'GAS FLOWRATE, lb/hr.:', T60, F12.3,/, 5X,'SUCTION PRESSURE, psia:', T60, F12.3,/, 5X,'DISCHARGE PRESSURE, psia:', T60, F12.3,/, 5X,'COMPRESSION RATIO:', T60, F12.3)</pre>
130 * *	<pre>WRITE (1, 130) T11, K, MW FORMAT (5X,'INLET TEMPERATURE, oF:',T60, F12.3,/, 5X,'RATIO OF SPECIFIC HEATS CAPACITIES, K:',T60,F12.3,/, 5X,'MOLECULAR WEIGHT OF GAS, lb/lb-mole:',T60, F12.3)</pre>
140 *	WRITE (1, 140) ZAV, AE FORMAT (5X,'AVERAGE COMPRESSIBILITY FACTOR, Z:',T60, F8.4,/, 5X,'ADIABATIC EFFICIENCY:',T60, F12.3)
150 * *	<pre>WRITE (1, 150) ADIAH, T22, HPADIA FORMAT (5X,'ADIABATIC HEAD, ft.:',T60, F12.3,/, 5X,'GAS DISCHARGE TEMPERATURE, oF:', T60, F12.3,/, 5X, 'GAS HORSEPOWER, hp:', T60, F12.3)</pre>
160	WRITE (1, 160) PROD FORMAT (5X,'MECHANICAL LOSSES, hp:',T60, F12.3)
170 *	WRITE (1, 170) BHPA, WORK FORMAT (5X,'TOTAL BRAKE HORSEPOWER, hp:',T60, F12.3,/, 5X,'WORK DONE, ft/lb-mole:',T60,F12.3)
180 *	WRITE (1, 180) AECAL FORMAT (5X,'CALCULATED ADIABATIC EFFICIENCY FROM',/, 5X,'POLYTROPIC EFFICIENCY:',T60, F12.3)
190	WRITE (1, 190) FORMAT (1H ,78(1H-))
с	FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR(12)
999	CLOSE (3,STATUS='KEEP') CLOSE (1) STOP END
с	***************************************

С THIS PROGRAM CALCULATES THE POLYTROPIC HEAD, THE ASSOCIATED GAS HORSEPOWER AND GAS DISCHARGE TEMPERATURE IN POLYTROPIC MACHINES С с ********* SUBROUTINE POLYTR REAL K, MW, N, MLOSS COMMON/PROG1/W, P1, P2, T11, T12, T21, T22, K, MW, N, ZAV, AE, AECAL, RC COMMON/PROG2/POLYH, ADIAH, HPPOLY, HPADIA, PE, PECAL, BHPP, BHPA COMMON/PROG3/WORK, PROD R = 1545.0T11 = T11 + 460.с CALCULATE THE COMPRESSION RATIO (DISCHARGE PRESS./SUCTION PRESS.) RC = P2/P1N = (K*PE)/(K*PE-K+1.0)С CALCULATE THE POLYTROPIC HEAD, POLYH: ft. VAL1 = (ZAV*R*T11)/MW VAL2 = (N/(N-1.0))*(RC**((N-1.0)/N)-1.0)POLYH = VAL1*VAL2 С CALCULATE THE GAS HORSEPOWER, HPPOLY: hp. HPPOLY = (W*POLYH)/(1.98*(10.0**6)*PE)C CALCULATE THE APPROXIMATE MECHANICAL LOSSES AS A PERCENTAGE OF GAS C POWER REQUIREMENT, PROD: hp. CALL MECHL (HPPOLY, MLOSS) C CALCULATE THE TOTAL BRAKE HORSEPOWER, BHPP: hp. BHPP = HPPOLY + PRODС CALCULATE THE GAS DISCHARGE TEMPERATURE, T12: OF. VAL3 = (POLYH*MW)/(ZAV*R)T12 = VAL3*((N-1.0)/N) + T11С CALCULATE THE WORK DONE 1b- mole/hr. ē CALCULATE THE BASIS IN moles. B = W/MWс CALCULATE THE WORK DONE WORK = (N/(N-1.0))*(R*T11*((P2/P1)**((N-1.0)/N)-1.0))WORK = (WORK*B)/(1.98*(10.0**6))T11 = T11 - 460.0T12 = T12 - 460.0RETURN END

С

с	**********************
c c c	THIS PROGRAM CALCULATES THE ADIABATIC HEAD, THE ASSOCIATED GAS HORSEPOWER AND GAS DISCHARGE TEMPERATURE
	SUBROUTINE ADIAB REAL K,MW,N,MLOSS COMMON/PROGI/W,P1,P2,T11,T12,T21,T22,K,MW,N,ZAV,AE,AECAL,RC COMMON/PROG2/POLYH,ADIAH,HPPOLY,HPADIA,PE,PECAL,BHPP,BHPA COMMON/PROG3/WORK, PROD
	PE = 0.75 R = 1545.0 T21 = T11 + 460.
с	CALCULATE THE COMPRESSION RATIO (DISCHARGE PRESS./SUCTION PRESS.)
	RC = P2/P1
с	CALCULATE THE ADIABATIC HEAD, ADIAH: ft.
	VAL4 = (ZAV*R*T21)/MW VAL5 = (K/(K-1.0))*(RC**((K-1.0)/K)-1.0) ADIAH = VAL4*VAL5
с	CALCULATE THE GAS HORSEPOWER, HPADIA: hp.
	HPADIA = (W*ADIAH)/(1.98*(10.0**6)*AE)
c c	CALCULATE THE APPROXIMATE MECHANICAL LOSSES AS A PERCENTAGE OF GAS POWER REQUIREMENT, PROD: hp.
	CALL MECHL (HPADIA, MLOSS)
с	CALCULATE THE TOTAL BRAKE HORSEPOWER, BHPA: hp.
	BHPA = HPADIA + PROD
с	CALCULATE THE GAS DISCHARGE TEMPERATURE, oF.
	T22 = T21*(RC**((K-1.0)/K)) T22 = T22 - 460.0
с	CALCULATE THE ADIABATIC EFFICIENCY FROM THE POLYTROPIC EFFICIENCY
	N = (K*PE)/(K*PE-K+1.0) VAL6 = (RC**((K-1.0)/K))-1.0 VAL7 = (RC**((N-1.0)/N))-1.0 AECAL = VAL6/VAL7
с	CALCULATE THE WORK DONE, hp
с	CALCULATE THE BASIS IN 1b-moles per hr.
	B = W/MW

C CALCULATE THE WORK DONE

```
WORK = (K/(K-1.0))*(R*T21)*((P2/P1)**((K-1.0)/K)-1.0)
WORK = (WORK*B)/(1.98*(10.0**6))
        RETURN
        END
с
с
с
с
с
        *******
                                                                *******
        THIS PROGRAM CALCULATES THE MECHANICAL LOSSES DUE TO
        FRICTION IN BEARINGS.
        SUBROUTINE MECHL (POLYL, MLOSS)
        COMMON/PROG3/WORK, PROD
        REAL MLOSS
        IF (POLYL .LT. 3000.0 ) THEN
        MLOSS = 0.03
        PROD = POLYL*MLOSS
        ELSEIF (POLYL .GT. 3000.0 .AND. POLYL .LT. 6000.0) THEN MLOSS\,=\,0.025
        PROD = POLYL*MLOSS
        ELSEIF (POLYL .GT. 6000.0 .AND. POLYL .LT. 10000.0) THEN
        MLOSS = 0.02
        PROD = POLYL*MLOSS
        ELSEIF (POLYL .GT. 10000.0) THEN MLOSS = 0.015
        PROD = POLYL*MLOSS
ENDIF
        RETURN
        END
```

PROGRAM PROG62

<u>_</u>			*****				
C							
с			M CALCULATES ALL PUMP HEAD CALCULATIONS REQUIRED TO WRITE				
с			IFICATION SUITABLE FOR A VENDOR INQUIRY OR PURCHASE				
с	ORDER.		PROGRAM FURTHER CALCULATES THE CENTRIFUGAL PUMP				
с			AND PUMP HORSEPOWER USING AN EQUATION DEVELOPED BY				
с	BY C. B	BY C. BRANAN (The Process Engineer's Pocket Handbook, Gulf Publishing					
с	Company	, но	uston 1976.)				
с	******	****	**********				
с							
с	FLUID	÷	FLUID NAME				
c	TEMP		FLUID OPERATING TEMPERATURE, OF.				
č	VIS	=					
č			FLOWRATE, gpm				
č	SPG	=	SPECIFIC GRAVITY OF LIQUID				
č	VP	_					
c	VP	-	VAPOR PRESSURE OF PUMPED FLUID AT PUMP SUCTION				
	VEAD		TEMPERATURE, psia.				
C	VESP		PRESSURE AT EQUIPMENT ON PUMP SUCTION, psia.				
с	SLL	=	SUCTION LINE LOSSES (PRESSURE DROP IN PIPING), psi.				
с	SSHD	=	SUCTION STATIC HEAD OF LIQUID ON PUMP SUCTION OR				
с			DISCHARGE, ft.				
с	NPSH	=	NET POSITIVE SUCTION HEAD ON PUMP, ft.				
с	DELP	=	PRESSURE AT EQUIPMENT ON PUMP DISCHARGE, psia.				
с	DSHD	=	DISCHARGE STATIC HEAD, ft.				
с	CVPD	÷	CONTROL VALVE PRESSURE DROP, psi.				
с	EXPD	*	HEAT EXCHANGER PRESSURE DROP, psi.				
c	ORPD	=	ORIFICE PRESURE DROP, psi.				
c	FURPD		FURNACE PRESSURE DROP, psi.				
č	DLL	-	DISCHARGE LINE LOSSES (PRESSURE DROP IN PIPING), psi.				
č	OTLD	×	OTHER LOSSES (DISCHARGE:-e.g. FURNACE), psi.				
č	OTLS	=	OTHER LOSSES (DISCHARGE. e.g. FORMACE), psi OTHER LOSSES (SUCTION:- e.g. STRAINER/FILTER), psi				
č	DELTAP						
č	DELPHT		PRESSURE DROP ACROSS PUMP, psi.				
			PRESSURE DROP ACROSS PUMP, ft.				
c	HBHP	=	HYDRAULIC BRAKE HORSEPOWER REQUIRED BY PUMP, bhp.				
с			(FOR HBHP CALCULATION, ASSUME PUMP EFFICIENCY=1.0)				
с	EFF	=					
с	F	=					
с		=					
с	BHP						
с		=	ft(0.433 x sp.gr.)				
с	ft	×	psi/(0.433 x sp.gr.)				
с	******	****	******************				
	CHARACT	FD+1					
			ORPD, OTLD, OTLS				
	KEAL N	ron,	ORFD, OILD, OILS				
			1/ FLUID, TEMP, VIS, SPG, VP				
	COMMON/DATA2/ GPM, VESP, SLL, SSHD, SSHDP						
	COMMON/DATA3/ DELP, DSHD, DSHDP, CVPD, EXPD						
	COMMON/DATA4/ ORPD, FURPD, DLL, OTLS, OTLD						
	COMMON /	DATA	5/ SP, SPGAGE, NPSH, DISP, DISPG				
	COMMON/DATA6/ DELTAP, DELPHT, HBHP, BHP, EFF						
			· · · · · · · · · · · · · · · · · · ·				

OPEN(UNIT=3,FILE='DATA62.DAT',STATUS='OLD',ERR=18) OPEN (UNIT =1, FILE='PRN')

5	READ (3, 5, ERR=19) FLUID FORMAT (A)
	READ (3, *, ERR=19) TEMP, VIS, SPG READ(3, *, ERR=19) VP, GPM READ(3, *, ERR=19) VESP, SLL, SSHD READ(3, *, ERR=19) DELP, DSHD,CVPD READ(3, *, ERR=19) EXPD, ORPD, FURPD READ(3, *, ERR=19) DLL, OTLS, OTLD
	GO TO 15
18 11	WRITE(*, 11) Format(6x,'file does not exist') Go to 999
19 12	WRITE(*, 12) Format(6x,'error message in the data value') Go to 999
15	CALL PUMPHA
10	WRITE(1, 10) FORMAT(26X,'CENTRIFUGAL PUMP SIZING',/,1H ,78(1H*))
20 * * * *	<pre>WRITE (1, 20) FLUID, TEMP, VIS, SPG, VP, GPM FORMAT (5X,'FLUID NAME:', T65, A,/,</pre>
30 * *	<pre>WRITE (1, 30) VESP, SLL, OTLS, SSHD FORMAT (5X,'PRESSURE AT EQUIPMENT, psia:', T65, F9.3,/, 5X,'SUCTION LINE LOSSES, psi:', T65, F9.3,/, 5X,'OTHER LOSSES (e.g. STRAINER/FILTER), psi:',T65,F9.3, /,5X,'SUCTION STATIC HEAD, ft.:', T65, F9.3)</pre>
40 * *	WRITE (1, 40) SSHDP, SPGAGE, SP FORMAT (5X,'SUCTION STATIC HEAD, psi:', T65, F9.3,/, 5X,'TOTAL PUMP SUCTION PRESSURE, psig:', T65, F9.3,/, 5X,'TOTAL PUMP SUCTION PRESSURE, psia:', T65, F9.3)
50	WRITE (1, 50) NPSH FORMAT (5X,'AVAILABLE NET POSITIVE SUCTION HEAD, ft.:', T65, F9.3)
60 * *	<pre>WRITE (1, 60) DELP, DSHD, DSHDP, CVPD FORMAT (5x,'DELIVERY PRESSURE, psia.:', T65, F9.3,/, 5x,'DISCHARGE STATIC HEAD, ft.:', T65, F9.3,/, 5x,'DISCHARGE STATIC HEAD, psi:', T65, F9.3,/, 5x,'CONTROL VALVE PRESSURE DROP, psi.:', T65, F9.3)</pre>
70 * *	<pre>WRITE (1, 70) EXPD, ORPD, FURPD, DLL, OTLD FORMAT (5X,'EXCHANGER PRESSURE DROP, psi.:', T65, F9.3,/, 5X,'ORIFICE PRESSURE DROP, psi.:', T65, F9.3,/, 5X,'FURNACE PRESSURE DROP, psi.:', T65, F9.3,/,</pre>

*	5X,'DISCHARGE LINE LOSSES, psi.:', T65, F9.3,/, 5X,'OTHER LOSSES, psi.:', T65, F9.3)
80 *	WRITE(1, 80)DISPG,DISP FORMAT(5X,'TOTAL PUMP DISCHARGE PRESSURE, psig:', T65, F9.3,/, 5X,'TOTAL PUMP DISCHARGE PRESSURE, psia:', T65, F9.3)
90 *	WRITE(1, 90)DELTAP,DELPHT FORMAT(5X,'TOTAL PUMP PRESSURE DROP, psi.:', T65, F9.3,/, 5X,'TOTAL PUMP PRESSURE DROP, ft.:', T65, F9.3)
100 *	WRITE(1, 100)HBHP FORMAT(5X,'HYDRAULIC BRAKE HORSEPOWER REQUIRED BY THE PUMP,' ' bhp:', T65, F9.3)
110 *	WRITE(1, 110)BHP FORMAT(5X,'ACTUAL BRAKE HORSEPOWER REQUIRED FOR PUMP OPERATION', ' bhp:', T65, F9.3)
120	WRITE (1, 120) EFF Format (5X,'PUMP EFFICIENCY, (%):', T65, F9.3)
130	WRITE (1, 130) FORMAT (1H ,78(1H-))
с	FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR(12)
	CLOSE(3,STATUS='KEEP') CLOSE (1)
999	STOP END
с с с с	**************************************
	SUBROUTINE PUMPHA
	CHARACTER*14 FLUID
	REAL NPSH, OTLS, OTLD, ORPD
	COMMON/DATA1/ FLUID, TEMP, VIS, SPG, VP COMMON/DATA2/ GPM, VESP, SLL, SSHD, SSHDP COMMON/DATA3/ DELP, DSHD, DSHDP, CVPD, EXPD COMMON/DATA4/ ORPD, FURPD, DLL, OTLS, OTLD COMMON/DATA5/ SP. SPGAGE, NPSH, DISP, DISPG

COMMON/DATA5/ SP, SPGAGE, NPSH, DISP, DISPG COMMON/DATA6/ DELTAP, DELPHT, HBHP, BHP, EFF

VAL1 = 0.4329*SPG SSHDP = SSHD*VAL1 SP = VESP + SSHDP- SLL- OTLS

```
SPGAGE = SP-14.7
с
        AVAILABLE NET POSITIVE SUCTION HEAD
        SLL1 = SLL/VAL1
        VESP1 = VESP/VAL1
        VP1 = VP/VAL1
DEF = VESP1-VP1
        NPSH = DEF + SSHD - SLL1
с
        PUMP DISCHARGE PRESSURE
        DSHDP = DSHD*VAL1
        DISP = DELP+DSHDP+CVPD+EXPD+ORPD+FURPD+DLL+OTLD
        DISPG = DISP-14.7
c
        TOTAL PUMP DELTAP = PUMP DISCH. PRES. - PUMP SUC. PRES.
        DELTAP = DISP-SP
        DELPHT = DELTAP/VAL1
        HYDRAULIC BRAKE HORSEPOWER REQUIRED BY PUMP
С
с
        PUMP EFFICIENCY IS ASSUMED = 1.0
        HBHP = (DELTAP*GPM)/(1715.0*1.0)
        THIS PROGRAM CALCULATES EFFICIENCY AND PUMP HORSEPOWER USING
с
        C. BRANAN'S EQUATION (e.g. C Brannan: The Process Engineer's Pocket Handbook, Gulf publishing Co.)
С
¢
        F = DELPHT
        G = GPM
        VAL2=0.2855*F
        VAL3=3.78*F*G*(10.0**(-4))
        VAL4=2.38*F*(G**2)*(10.0**(-7))
        VAL5=5.39*(F**2)*(10.0**(-4))
        VAL6=6.39*(F**2)*G*(10.0**(-7))
        VAL7=4.0*(F**2)*(G**2)*(10.0**(-10))
        EFF=(80.0-VAL2+VAL3-VAL4+VAL5-VAL6+VAL7)
        BHP=(100.0*HBHP)/EFF
```

RETURN END

PROGRAM PROG63

```
c
С
        THIS PROGRAM WILL CALCULATE THE PUMP EFFICIENCY BASED ON
00000000000
        THE FLUID FLOW RATE AND THE HEAD USING THE EQUATIONS
        DEVELOPED BY C. BRANAN (The Process Engineer's Pocket Handbook,
        Gulf Publishing Company, Houston 1976).
        н
                =
                     HEAD, ft.
                     VOLUMETRIC FLOWRATE, gpm.
                =
        ٥
        HBHP
                     HYDRAULIC HORSEPOWER, bhp.
                =
                     BRAKE HORSEPOWER, bhp.
        BHP
                =
         DIMENSION EFF(20), H(20), Q(20), HBHP(20), BHP(20)
        OPEN (UNIT=3, FILE='DATA63.DAT', STATUS='OLD', ERR=18)
OPEN (UNIT=1, FILE='PRN')
        READ (3, *, ERR=19) N
        DO 10 I=1, N
READ (3, *, ERR=19) Q(I), H(I)
 10
        CONTINUE
        GO TO 44
        WRITE (*, 20)
FORMAT (6X, 'FILE DOES NOT EXIST')
 18
 20
        GO TO 999
        WRITE (*, 21)
FORMAT (6X, 'ERROR MESSAGE IN THE DATA VALUE')
 19
 21
        WRITE (1, 90)
FORMAT (///,25X,'EFFICIENCY AND BRAKE HORSEPOWER CALCULATIONS',/
 44
 90
                 1H ,78(1H*),//)
     .
       WRITE(1, 95)
FORMAT (5X,'FLOWRATE gpm', 10X,'HEAD ft.', 6X,'EFFICIENCY %',
6X,'BRAKE HORSEPOWER Bhp',/1H , 78(1H*))
 95
     *
        SPGR=1.0
        DO 35 I=1,N
        IF (Q(1), EQ. 0.0) THEN
EFF(1) = 0.0
        HBHP(1) = 0.0
        ELSE
        ENDIF
```

	VAL1 = 0.2855*H(I) VAL2 = 3.78*H(I)*Q(I)*(10.0**(-4)) VAL3 = 2.38*H(I)*(Q(I)**2)*(10.0**(-7)) VAL4 = 5.39*(H(I)**2)*(10.0**(-4)) VAL5 = 6.39*(H(I)**2)*Q(I)*(10.0**(-7)) VAL6 = 4.0*(H(I)**2)*(Q(I)**2)*(10.0**(-10)) EFF(I)=(80.0-VAL1+VAL2-VAL3+VAL4-VAL5+VAL6)
с с	HYDRAULIC BRAKE HORSEPOWER REQUIRED BY THE PUMP PUMP EFFICIENCY IS ASSUMED = 1.0
	HBHP(I)=(Q(I)*H(I)*SPGR)/(3960.0*1.0) BHP(I)=HBHP(I)/(EFF(I)/100.0)
35	CONTINUE
	DO I =1, N
96	WRITE (1, 96)Q(I),H(I),EFF(I),BHP(I) FORMAT (5X, F8.1, 10x, F8.1, 10X, F8.1, 10X, F8.1)
	ENDDO
100 *	<pre>WRITE(*, 100) FORMAT (5X , 'FLOWRATE gpm', 10X, 'HEAD ft.', 6X 'EFFICIENCY %', 6X,'BRAKE HORSEPOWER Bhp',/1H ,78(1H*))</pre>
110 45	DO 45 I =1, N WRITE(*, 110)Q(I), H(I), EFF(I), BHP(I) FORMAT (5X, F8.1, 10X, F8.1, 10X, F8.2, 10X, F8.1) CONTINUE
120	WRITE (1,120) FORMAT (1H ,78(1H-))
с	FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR(12)
	CLOSE (3, STATUS='KEEP') CLOSE (1)
9 9 9	STOP END

CHAPTER 7

Mass Transfer

INTRODUCTION

Unit operation techniques in the chemical process industries (CPI) effect the separation of a mixture or mixtures into their components. These operations are called diffusional or mass transfer operations. Techniques employed in the CPI to effect these separations are: distillation, absorption, liquid extraction, drying, leaching, crystallization, and gas adsorption. These processes account for 40% to 70% of both the capital and operating costs in the CPI. Separation operations significantly affect energy consumption, product costs, and manufacturing profits. About 43% of the energy consumed by the U.S. chemical process industries is used for separation processes [1]. The method depends on the state (such as liquid, solid, or gaseous) of the components.

The engineer must adequately size equipment for separating and recovering products from multicomponent feed streams. In addition, he or she should consider whether the design is either stagewise or continuous, tray or packed. The trays act as individual stages and produce stepwise changes in concentration, while packed towers give gradual changes in concentration. Designing distillation equipment requires knowledge of vapor-liquid equilibria in multi-component mixtures.

This chapter considers the vapor-liquid equilibrium of mixtures, conditions for bubble and dew points of gaseous mixtures, isothermal equilibrium flash calculations, the design of distillation towers with valve trays, packed tower design, Smoker's equation for estimating the number of plates in a binary mixture, and finally, the computation of multicomponent recovery and minimum trays in distillation columns.

VAPOR-LIQUID EQUILIBRIUM

A common operation in the CPI for the separation of fluid mixtures into their components is distillation. The distillation process is a separation technique that depends on the difference between the compositions of liquid and vapor phases at equilibrium. This implies that the temperatures and pressures of the phases must be the same, and no composition change occurs with time. Equilibrium can be achieved after a long period of thorough mixing and contact between the two phases.

An ideal gas is, by definition, one that follows the ideal gas laws:

$$PV = nRT$$
(7-1)

where P = absolute pressure

V = total volume

n = number of moles of gas

R = ideal gas constant

T = absolute temperature

The conditions under which a given component or mixture approaches ideal behavior depend on the critical temperature and critical pressure. Other principles of gas behavior are Dalton's law of additive pressures and Amagat's law of additive volumes. These are:

$$\pi = p_A + p_B + \dots \tag{7-2}$$

and

$$V_{\rm T} = V_{\rm A} + V_{\rm B} + \dots$$
 (7-3)

where π = total system pressure

p = partial pressure $V_{A} = volume of component A$ $V_{B} = volume of component B$ $V_{t} = total volume of system$ A,B = components

These laws are correct when conditions are such that each component and the mixture obey the ideal gas law. The equilibrium between two phases can be related to the equality of the chemical potential and defined in terms of the Gibbs free energy as

$$\mu_{i} = \left[\frac{\partial G}{\partial \mu_{i}}\right]_{T,P,n_{j}} = \overline{G}_{i}$$
(7-4)

where \overline{G}_i is the partial molar Gibbs free energy. From the thermodynamic relationship between Gibbs free energy, temperature and pressure, and at constant temperature,

$$dG = VdP \tag{7-5}$$

where G is the molar Gibbs free energy and V is the molar volume.

Considering a pure fluid of component i and substituting Equation 7-1 into Equation 7-5,

$$dG = RT \frac{dP}{p}$$
(7-6)
= RTd(ln P)

The free energy from Equation 7-6 is restricted to ideal gases, but can be modified to include real fluids by introducing a new function defined as the fugacity. Equation 7-6 becomes

$$dG_i = RTd(\ln f_i) \tag{7-7}$$

where f_i is the fugacity of pure i with units of pressure. The fugacity becomes equal to the pressure as the pressure approaches zero.

Thus,

$$\lim \frac{f_i}{P} = 1.0 \tag{7-8}$$

$$P \to 0$$

For a single component in a mixture at constant temperature, the fugacity is defined by

$$d\overline{G}_{i} = RTd(\ln \overline{f}_{i})$$
(7-9)

and

$$\lim \frac{\bar{f}_i}{y_i P} = 1.0 \tag{7-10}$$
$$P \to 0$$

If a liquid mixture at temperature T and pressure P is in equilibrium with a vapor mixture at the same temperature and pressure, therefore at an equilibrium condition, the thermodynamic criterion will be 472 Fortran Programs for Chemical Process Design

$$\mathbf{f}_{i}^{\mathrm{V}} = \mathbf{f}_{i}^{\mathrm{L}} \tag{7-11}$$

where $f_i^v =$ fugacity of component i in the vapor phase $f_i^{L} =$ fugacity of component i in the liquid phase

The fugacity of a component in a mixture depends on the temperature, pressure, and composition of that mixture. The fugacity, f_{i}^{L} , of any component, i, in the liquid phase is related to the composition (mole fraction) of the component in that phase by

$$\mathbf{f}_{i}^{L} = \boldsymbol{\gamma}_{i} \mathbf{x}_{i} \mathbf{f}_{i}^{oL} \tag{7-12}$$

where γ_i = the activity coefficient of component i

 $x_i = mole fraction of component i$ $<math>f_i^{oL} = the fugacity of component i in the reference (or standard)$ state

The fugacity of pure liquid **i** at temperature T and pressure P is given by

$$f_i^{oL} = P_{vp,i}(T)\phi_i^s(T)\exp\left[\int_{P_{vp,i}}^{P} \frac{V_i^L(T,P)}{RT}dP\right]$$
(7-13)

where $P_{vp,i}$ = the vapor pressure of component i φ_i^s = the vapor phase fugacity coefficient of pure saturated vapor of component i

The exponential term in Equation 7-13 is a correction factor for the effect of pressure on liquid-phase fugacity and is known as the Poynting factor. In Equation 7-13, V_i^L can be replaced by the partial molar volume of component i in the liquid solution for greater accuracy. For low to moderate pressure, V_i^L is assumed as the saturated liquid molar volume at the specified temperature. Equation 7-13 is simplified to give

$$f_i^{oL} = P_{vp,i} \phi_i^s \exp\left[\frac{V_i^{Ls}(P - P_{vp,i})}{RT}\right]$$
(7-14)

Equation 7-14 is used to calculate the reference state fugacity of liquids. Any equation of state can be used to evaluate ϕ_i^s . For low to moderate pressures, the virial equation is the simplest to use. The fugacities of pure gases and gas mixtures are needed for estimating many thermodynamic properties, such as entropy, enthalpy, and fluid phase equilibria. For pure gases, the fugacity is

$$\ln f = \frac{1}{RT} \int_0^P V dP \tag{7-15}$$

The fugacity can be calculated from Equation 7-15 once the P-V-T behavior of the fluid is known from an equation of state. The fugacity coefficient is the ratio of the fugacity of a substance to its pressure. For a pure substance,

$$\phi = \frac{f}{P} \tag{7-16}$$

For the vapor phase, the composition is nearly always expressed by the mole fraction, y_i

$$\phi_i = \frac{f_i^{\nu}}{y_i P} \tag{7-17}$$

For a mixture of ideal gases, $\phi_i = 1$. The fugacity coefficient, ϕ_i , depends on temperature and pressure. The fugacity coefficient is normalized, such that as $P \rightarrow 0$, $\phi_i \rightarrow 1$ for all components. At a low pressure, ϕ_i is assumed to be unity.

When the volume is known as a function of the pressure, either from direct measurement or from an empirical equation of state, changes in fugacity may be found by integration. Thus

$$\ln \frac{f_2}{f_1} = \frac{1}{RT} \int_{P_1}^{P_2} V dP$$
 (7-18)

Since $\phi = 1$ when P = 0, absolute values of the fugacity coefficient can be determined. Therefore,

$$\ln \phi = \ln \frac{f}{P} = \frac{1}{RT} \int_0^P \left(V - \frac{RT}{P} \right) dP$$

$$= \int_0^P (Z - 1) \frac{dP}{P}$$
(7-19)

Methods for estimating fugacities of liquid and vapor phases are given by Reid, *et al.* [2], and by Walas [3].

Raoult's law states that the partial pressure in the liquid phase can be expressed as

$$\mathbf{p}_{\mathbf{A}} = \mathbf{x}_{\mathbf{A}} \mathbf{p}_{\mathbf{A}}^{\circ} \tag{7-20}$$

where x_A is the mole fraction of component A, and p_A^o is the vapor pressure of component A.

For the vapor phase, Dalton's law can be expressed as:

$$p_A = \pi y_A \tag{7-21}$$

where y_A is the mole fraction of component A. Combining Equation 7-20 and Equation 7-21

$$x_{A}p_{A}^{\circ} = \pi y_{A}$$

$$K = \frac{y_{A}}{x_{A}} = \frac{p_{A}^{\circ}}{\pi}$$
(7-22)

where K is the equilibrium constant.

Few multicomponent systems exist for which completely generalized equilibrium data are available. The most widely available data are those for vapor-liquid systems, and these are frequently referred to as vapor-liquid equilibrium distribution coefficients or K value. The K values vary with temperature and pressure, and a selectivity that is equal to the ratio of the K values is used. For vapor-liquid systems, this is referred to as the relative volatility and is expressed for a binary system as

$$\alpha = \frac{K_A}{K_B}$$
(7-23)

where

$$\mathbf{K}_{\mathrm{A}} = \mathbf{y}_{\mathrm{A}} / \mathbf{x}_{\mathrm{A}}$$

and

* *

$$\kappa_{\rm B} = y_{\rm B} / x_{\rm B}$$

$$\alpha_{\rm A-B} = \frac{y_{\rm A} \cdot x_{\rm B}}{x_{\rm A} \cdot y_{\rm B}}$$
(7-24)

Equation 7-24 can be expressed in terms of the more volatile component A, as

$$\mathbf{y}_{\mathrm{A}} = \frac{\alpha \mathbf{x}_{\mathrm{A}}}{1 + (\alpha - 1)\mathbf{x}_{\mathrm{A}}} \tag{7-25}$$

where

 $x_{A} + x_{B} = 1$

and

 $y_A + y_B = 1$

Equation 7-25 relates equilibrium compositions x_A and y_A in terms of the relative volatility. If the assumption is that α is independent of temperature and composition, then Equation 7-25 becomes the equation of the equilibrium line or the x-y curve, as shown in Figure 7-1. Figure 7-1 can be used to determine the compositions of vapor and liquid in equilibrium.

BUBBLE POINT OF GAS MIXTURES

An equilibrium between a liquid and a vapor implies that the liquid is on the point of boiling, is producing bubbles of vapor within it, and the

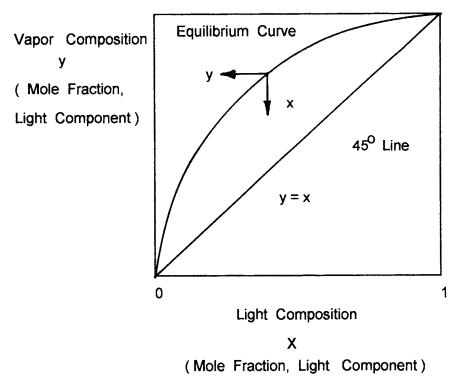


Figure 7-1. X-Y diagram for a binary system.

vapor is on the point of condensing or forming a dew. These conditions are referred to as saturated liquid and saturated vapor. For the particular total pressure, the liquid is at its bubble point temperature and the vapor at its dew point temperature. The amount of vapor formed is assumed to be so small as to have no effect on the liquid composition. For a binary system,

$$\mathbf{x}_1 = \frac{1 - \mathbf{K}_2}{\mathbf{K}_1 - \mathbf{K}_2} \tag{7-26}$$

and

$$\mathbf{x}_2 = 1.0 - \mathbf{x}_1 \tag{7-27}$$

In many cases where there are more than two components in the liquid mixture, no direct solution as Equation 7-26 exists. The basic definition of the equilibrium coefficient is

$$\mathbf{y}_{i} = \mathbf{K}_{i}\mathbf{x}_{i} \tag{7-28}$$

For a stable system, the total mole fraction is unity, which is

$$\sum \mathbf{y}_{i} = 1.0 = \sum \mathbf{K}_{i} \mathbf{x}_{i}$$
(7-29)

Equation 7-29 is used to determine the bubble point temperature and pressure. In using Equation 7-29, the temperature or pressure is fixed, while the other parameter is varied until the criterion for a stable system is satisfied. A combination of temperature and pressure is altered, if the summation of the calculated vapor composition is different from unity. There is no known direct method that will allow a reasonable estimation of the amount of change required. However, Dodge [4], Hines and Maddox [5] have provided techniques for reducing the number of trials that is required.

DEW POINT CALCULATIONS

The dew point of a vapor is that combination of temperature and pressure at which the first drop of liquid condenses. The dew point criterion can be expressed as:

$$\sum x_{i} = 1.0 = \sum \frac{y_{i}}{K_{i}}$$
(7-30)

For binary mixtures of components 1 and 2, then

$$\mathbf{y}_{i} = \frac{\mathbf{K}_{1}(1 - \mathbf{K}_{2})}{\mathbf{K}_{1} - \mathbf{K}_{2}} \tag{7-31}$$

and

$$y_2 = 1.0 - y_1 \tag{7-32}$$

These two equations can be used to calculate the composition of a binary vapor that will begin to condense at a given temperature and pressure.

EQUILIBRIUM FLASH COMPUTATIONS

Flash vaporization calculations involving multicomponent mixtures are essential to the design and successful operation of many processes. These calculations are often required to determine the condition of the feed to a fractionating column or to determine the flow of vapor from a reboiler or condenser.

Flash calculations often involve trial-and-error solutions that are time consuming, tedious, and subject to error if performed manually. Coker [6] has developed an improved iterative convergence method first suggested by Oliver [7] and later modified by Kostecke [8], for isothermal equilibrium flash computations. Multicomponent flash (isothermal and adiabatic) computations are incorporated as part of overall process simulation and equipment design. However, single-stage flash fractionation processes are also employed to obtain a separation of the light components in a feed and as a preliminary step before a multicomponent fractionation column, such as crude oil distillation. Table 7-1 lists ways of producing two-phase mixtures from a single phase at appropriate conditions. The last process in the table typifies the well head separation that takes place in an oil field. It is known as a flash process because the

Initial	Action to Produce Two-Phase Mixture
Gas	Cool, possibly after initial compression
Gas	Expand through a valve or an engine
Liquid	Heat to achieve partial vaporization
Liquid	Reduce pressure through a valve, if close to saturation

Table 7-1 Ways to Produce Two-Phase Mixture

vapor forms due to the rapid drop in the pressure.

Fundamentals

To carry out an appropriate flash calculation, the pressure, P, and the temperature, T, must be known. If the values of P and T in the separating vessel are fixed, the value of P must not be so high that the two phases cannot exist at any value of T. Nor must T lie outside the bubble point and dew point range corresponding to P. For a valid two-phase equilibrium calculation, the following relationship must be satisfied:

$$T_{bp} < T_s < T_{dp} \tag{7-33}$$

where T_{bp} = the bubble point temperature

 $T_s =$ the specified temperature

 T_{dp} = the dew point temperature

The existence of a valid two-phase flash can be verified with the design equations for the bubble point, dew point, and equilibrium data calculated at the specified pressure and temperature. The design equations for the bubble and dew points are:

Bubble point

$$f_{1} = \sum_{i=1}^{n} K_{i} X_{i}$$
(7-34)

Dew point

$$f_{2} = \sum_{i=1}^{n} Y_{i} / K_{i}$$
(7-35)

Table 7-2 illustrates the phase condition using the liquid-vapor data associated with the specified pressure and temperature.

From Table 7-2, the conditions for a valid two-phase equilibrium flash are:

$$f_{1} = \sum_{i=1}^{n} K_{i} n_{i} > 1.0$$
(7-36)

and

$$f_2 = \sum_{i=1}^{n} n_i / K_i > 1.0$$
(7-37)

- <u></u>	$\mathbf{f}_1 = \sum_{i=1}^n \mathbf{K}_i \mathbf{n}_i$	$f_2 = \sum_{i=1}^n n_i / K_i$
Subcooled liquid	<1	>1
Bubble point	=1	>1
Two-phase condition	>1	>1
Dew point	>1	=1
Superheated vapor	>1	<1

Table 7-2 Equilibrium Flash Criteria

Feed analyses of component concentrations are normally not available for complex hydrocarbon mixtures with a final boiling point of more than about 38°C (e.g. n-pentane), unless such a feed is broken down into pseudo-components (narrow boiling fractions). This enables the mole fraction and equilibrium constant, K, to be estimated, and consequently, flash calculation of the mixture can be carried out. A source of K values for light hydrocarbon systems is the De-Priester [9] charts, which are shown in Figures 7-2 and 7-3. These charts give the K values over a wide range of temperature and pressure. Because the charts are nomographs, a straight edge connecting the temperature and pressure for which K values are required will intersect curves for each compound at its K value. These K values have been obtained by calculating fugacities from an equation of state. Use of these charts for exact determination of K values requires trial-and-error calculations. Hadden and Grayson [10] have presented correlations for hydrocarbon vapor-liquid distribution ratios, and the charts from their work are shown in Figures 7-4 and 7-5. These charts can be readily used for determining the vaporliquid distribution coefficients. Figure 7-6 shows a continuous equilibrium flash fractionation process.

The Equations

The following equations are used for the multicomponent equilibrium flash calculations.

$$f_1 = \sum_{i=1}^{n} n_i K_i < 1 \quad \text{all liquid}$$
(7-38)

(text continued on page 484)

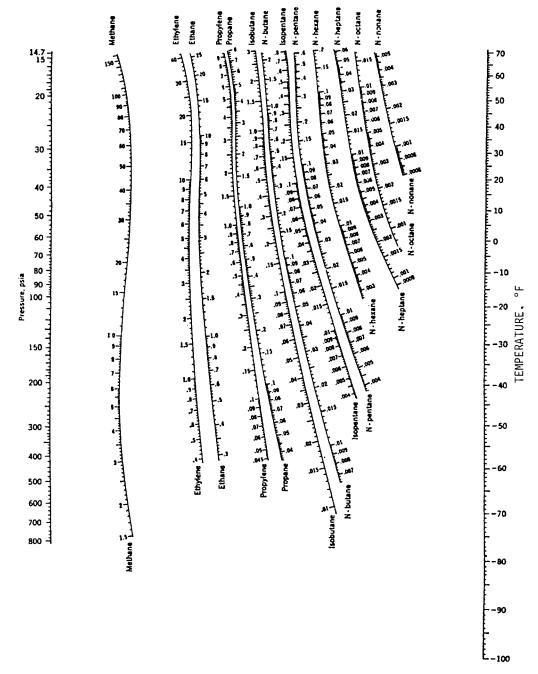


Figure 7-2. K-values for hydrocarbons, low temperature. Source: De-Priester [9].

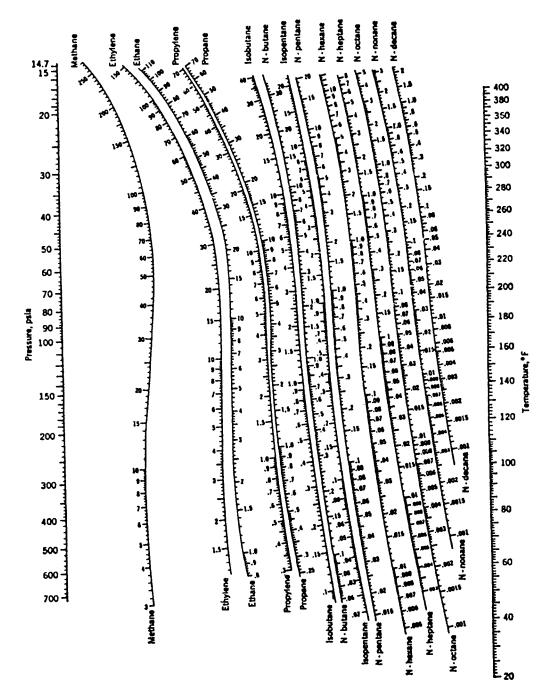
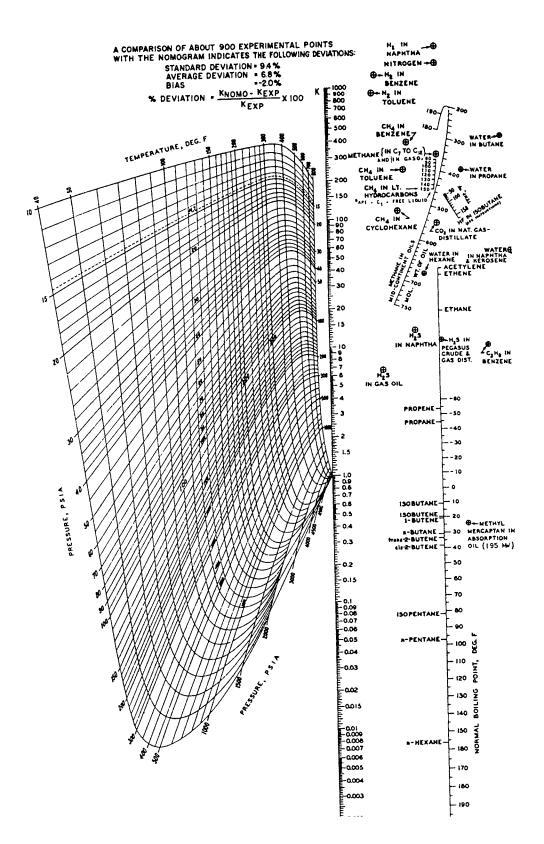


Figure 7-3. K-values for hydrocarbons, high temperature. Source: De-Priester [9].



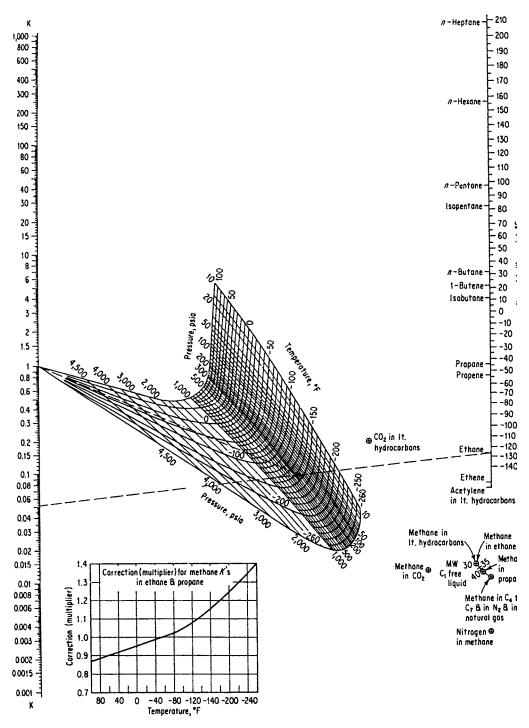


Figure 7-5. Vapor-liquid equilibrium constants, -260 to +100°F. Source: Hadden and Grayson [10].

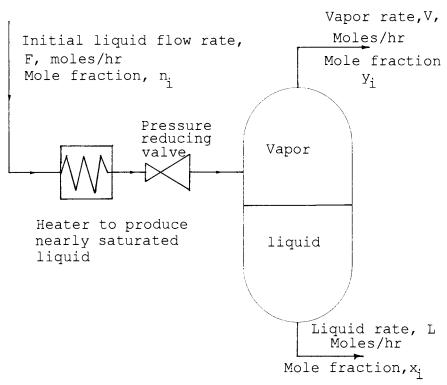


Figure 7-6. Continuous equilibrium flash fractionation.

(text continued from page 479)

$$f_{2} = \sum_{i=1}^{n} n_{i}/K_{i} < 1 \quad \text{all vapor}$$

$$C_{i} = \frac{\left[M_{i}(K_{i}-1)(R+1)\right]}{(K_{i}+R)}$$
(7-39)
(7-40)

where R = the liquid-vapor (L/V) ratio. The new L/V ratio for each iterative calculation, R', is determined from:

$$R' = \frac{[FR - E(R+1)]}{[F + E(R+1)]}$$
(7-41)

The constants E and F are determined by

$$E = \sum_{i=1}^{n} C_i$$
(7-42)

$$F = \sum_{i=1}^{n} (C_i)^2 / M_i$$
 (7-43)

The Algorithm

The computation then determines whether |E| < 0.001. If |E| is not less than 0.001, R is set equal to R' and the calculations of Equations 7-40, 7-41, 7-42 and 7-43 are repeated until |E| < 0.001.

Once |E| < 0.001, the calculation proceeds to find the total moles of component i in both the liquid and the vapor phases, that is, L_i and V_i . These are given by

$$L_{i} = \frac{M_{i}R}{(K_{i} + R)}$$
(7-44)

and

$$\mathbf{V}_{i} = \mathbf{M}_{i} - \mathbf{L}_{i} \tag{7-45}$$

The mole fractions of components in the feed, the liquid and vapor phases are evaluated as follows:

$$n_{i} = \frac{M_{i}}{\sum_{i=1}^{n} M_{i}}$$
(7-46)

$$X_{i} = \frac{L_{i}}{\sum_{i=1}^{n} L_{i}}$$
(7-47)

and

$$\mathbf{Y}_{i} = \frac{\mathbf{V}_{i}}{\sum_{i=1}^{n} \mathbf{V}_{i}}$$
(7-48)

where

$$\sum_{i=1}^{n} n_{i} = \sum_{i=1}^{n} X_{i} = \sum_{i=1}^{n} Y_{i} = 1$$
(7-49)

The method described here is based on the vapor-liquid equilibrium relationships given in handbooks available from the Gas Processors Suppliers Association. This technique will handle flash calculations with feed streams containing up to 15 components. As an added feature, the calculation will check the feed composition at flash conditions for dew point or bubble point condition (i.e., whether the feed is either all vapor or all liquid). These checks are done before the flash calculations are started. If the feed is above the dew point or below the bubble point, an appropriate message is displayed on the computer screen. A default value for R (L/V) = 1.0 is used to start the iterative process.

Nomenclature

E,F = constants

- K_i = equilibrium flash constant for each component in the feed stream
- L_i = total moles of component i in the liquid phase
- L = total moles of liquid at equilibrium conditions
- M_i = total moles of component i in the feed stream
- n_i = mole fraction of component i in the feed stream
- R = liquid-vapor (L/V) ratio
- R' = new L/V ratio for each iteration calculation
- V = total moles of vapor at equilibrium conditions
- V_i = total moles of component i in the vapor phase
- X_i = mole fraction of component i in the liquid phase
- Y_i = mole fraction of component i in the vapor phase

TOWER SIZING FOR VALVE TRAYS

Introduction

Many types of trays are used in both fractionating and absorption columns. In a fractionating column, the bubble caps with the weirs and downcomers maintain a liquid level on the trays. The liquid flows across the tray via the downcomer and across the next tray in the opposite direction, and the vapor flows up through the caps and the slots, thus mixing with the liquid. Figures 7-7 and 7-8 show a section of the column distributor and different types of packings. Figure 7-9 illustrates the vapor flows through bubble cap, sieve, and valve trays. The riser in the bubble cap can aid the design of the tray to prevent liquid from "weeping" through the vapor passage. Sieve or valve trays control weeping by vapor velocity. The bubble cap tray has the highest turn down

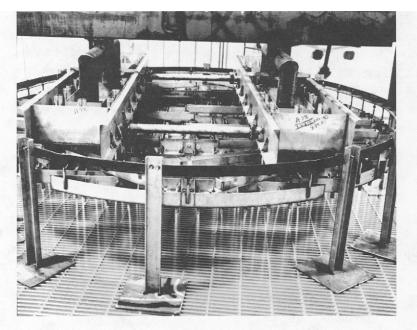
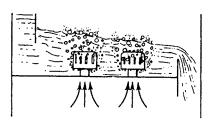
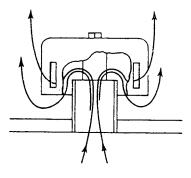


Figure 7-7. Nutter Engineering MTS-109 multi-pan two-stage liquid distributor Courtesy of Nutter Engineering.

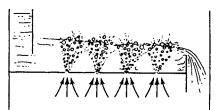


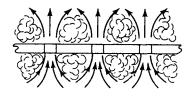
Figure 7-8. Nutter ring random packing. Courtesy of Nutter Engineering.





(o)





(b)

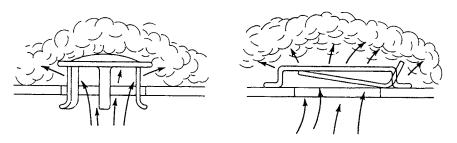


Figure 7-9. Flow through vapor passages. (a) Vapor flow through bubble cap. (b) Vapor flow through perforations. (c) Vapor flow through valves. Source: Van-Winkle [11].

ratio with the design of 8:1 to 10:1 ratio. Such trays are commonly used in glycol dehydrating columns [11].

Valve and sieve trays are favored because of lower cost and increased capacity over bubble cap trays for a given tower diameter. Sieves or perforated trays are plates with holes for vapor passage. It is simple to construct and the least expensive of the three designs. Generally, the sieve tray has a higher capacity. At low vapor rates, it is susceptible to "weeping" or dumping of the liquid through the holes. Furthermore, sieve trays are very good in fouling applications or when solids are present. They can have large holes because they are highly resistant to clogging and are easy to clean. In addition, their turndown ratio is limited.

Trayed columns give satisfactory operation over a wide range of liquid and vapor loadings. Correspondingly, valve trays can maintain high efficiencies over a wider range of operating liquid and gas rates than with sieve trays. Valve units are more complex mechanically than sieve trays, and therefore are more expensive to fabricate. The liquid and vapor rates can vary independently over a broad range in the column resulting in a satisfactory operation. At low vapor rates, unsatisfactory tray dynamics may be characterized by dumping of liquid, uneven distribution, and vapor pulsation. Alternatively, at high vapor rates, the tower floods, as the liquid is backed-up in the downcomers. At low liquid rates, poor vapor-liquid contact can result. Correspondingly, at high liquid rates, flooding and dumping can also result as the liquid capacities are exceeded in the downcomers. Valve travs are designed to have better turndown properties than sieve trays, and therefore are more flexible for varying feed rate. At the design vapor rate, valve trays have about the same efficiency as sieve trays. They can be designed for a lower pressure drop (ΔP) than sieve trays. However, they are susceptible to fouling or plugging, if dirty solutions are distilled.

Here, valve trays are used for sizing the columns. The procedure uses the Tower sizing graphs and tables from Nutter [12] and Blackwell's [13] correlation.

The Equations

The following expressions are used for the design of valve trays columns:

The density radical, R_D , is defined by

$$R_{\rm D} = \left(\frac{\rho_{\rm V}}{\rho_{\rm V} - \rho_{\rm L}}\right)^{0.5} \tag{7-50}$$

The velocity at zero liquid load is:

$$V = \exp \left\{ A + B(\ln X) - C(\ln X)^{2} + D(\ln X)^{3} - E(\ln X)^{4} \right\}$$
(7-51)

where

$$X = \frac{\sigma}{\rho_{\rm v}} \tag{7-52}$$

A, B, C, D, and E are constants:

A = 0.22982 B = 0.46605 C = 0.03452 D = 0.00415E = 0.00017

Equation 7-51 is valid for $0.1 \le X \le 3000$.

The tray spacing factor, T_{SF} , is defined by

$$T_{SF} = G + H(1nX) \tag{7-53}$$

where Equation 7-53 is valid for $0.1 \le X \le 5000$. The constants G and H for different tray spacing are given in Table 7-3. Figure 7-10 illustrates the plots of T_{sF} versus $X(\sigma/\rho_v)$ for different tray spacing.

The operating volumetric vapor flow rate, Q_v , is determined by

$$Q_{v} = \frac{W_{v}}{(3600)(\rho_{v})}, \quad ft^{3}/s$$
(7-54)

The operating volumetric liquid flow rate, Q_L, is determined by

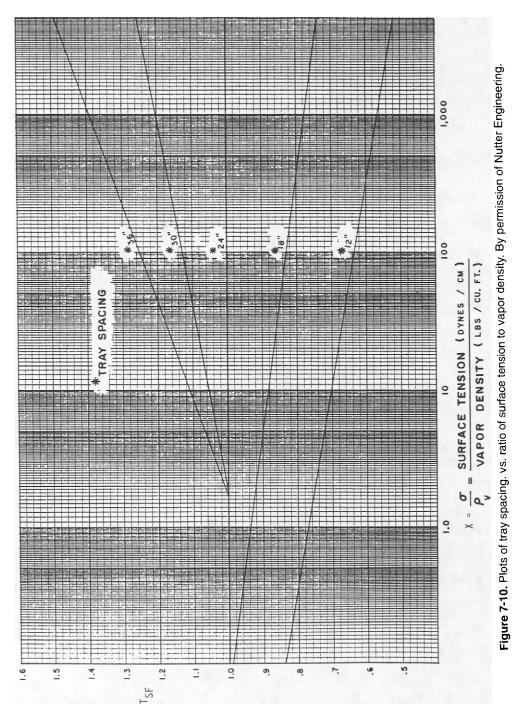
$$Q_{L} = \frac{W_{L}}{(3600)(\rho_{L})}, \quad ft^{3}/s$$
(7-55)

The bubbling area, A_b , is given by

Tray Spacing (inches)	G	Н	
12	0.77174	-0.02964	
18	0.93655	-0.02310	
24	$T_{sF} = 1.0$ for all values of X		
30	0.98057	0.03220	
	$T_{SF} = 1.0$ when X	< 1.85	
36	0.96583	0.06162	
	$T_{SF} = 1.0$ when X	K < 1.85	

 Table 7-3

 Values of G and H for Each Tray Spacing





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$$A_{b} = \frac{(Q_{v})(R_{D}) + Q_{L}}{(T_{sF})(V)(R_{D})(Foam fac)}, \quad ft^{2}$$
(7-56)

If $(V)(R_D) > 0.5$, then replace 0.5 for $(V)(R_D)$ in Equation 7-56. Table 7-4 gives recommended values of the foam factor and residence time.

Using Table 7-4 to obtain both the foam factor (Foam fac) and the downcomer residence time, τ , the downcomer area, A_{DC} , is determined by

$$A_{DC} = \frac{(\tau)(Q_{L})(12)}{(Foam fac)(T_{s})}, \quad ft^{2}$$
(7-57)

The expression for the safety factor, S_{Fac} , is given by

$$S_{Fac} = K(X)^{M}$$
(7-58)

Equation 7-58 is valid for $3.0 \le X \le 50,000$.

where K = 0.91146M = -0.03821

The tower area, A_{T} , is determined by

$$A_{T} = \frac{2(A_{DC}) + A_{b}}{S_{Fac}}, \quad ft^{2}$$
 (7-59)

Table 7-5 shows the tower diameter versus its cross-sectional areas, and Tables 7-6 to 7-9 give the recommended downcomer area and bubbling area distribution ratios.

The column diameter, D, is given by

$$D = \left(\frac{4.A_{T}}{\pi}\right)^{0.5}$$
(7-60)

The number of tray passes required is determined from Table 7-10.

Figures 7-11 and 7-12 show the dimensions of the various downcomers and the distribution area for the number of passes.

Nomenclature

 A_b = bubbling area, ft² A_{DC} = downcomer area, ft² A_T = tower area, ft²

Fractionators	Foam factor	Downcomer Res. Time, Sec.
Straight Run	· · · · · · · · · · · · · · · · · · ·	
Hydrocarbon	1.0	4.0
Low MW Alcohols	1.0	3.5
Crude Fractionator	1.0	4.5
Special		
Rich Oil (Top) Demeth. or Deeth.	0.85	4.5
Rich Oil (BTM) Demeth. or Deeth.	1.0	4.5
Hydrocarbon Still (Top)	1.0	4.0
Hydrocarbon Still (BTM)	1.0	4.5
MEA-DEA Still	0.85	4.5
Glycol-DGA Still	0.8	4.5
Sulfinol Still	1.0	5.0
H ₂ S Stripper	0.9	4.0
Sour Water Stripper	0.5-0.7	4.5
O ₂ Stripper	1.0	3.0
Refrigerated Demeth. or Deeth. (Top)	0.8	8.0
Refrigerated Demeth. or Deeth. (BTM)	1.0	5.0
Absorbers		
Oil-Ambient Temp. (Above 0°F)	0.85	4.5
Oil-Low Temp. (Below 0°F)	0.95	4.5
DGA-DEA-MEA (Contactor)	0.75	4.5
Glycol Contactor	0.65	5.0
Sulfinol Contactor	1.0	5.0
Vacuum Towers		
Crude Vacuum	1.0	5.0

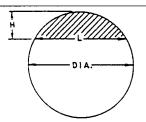
 Table 7-4

 Recommended Values for Foam Factor and Residence Time

Tower Diameter		Area	Tower D	Diameter	Area
Feet	Inches	Sq. Ft.	Feet	Inches	Sq. Ft.
2'-0''	24"	3.142	21'-0"	252"	346.361
2'-6''	30"	4.909	21'-6"	258"	363.050
30"	36"	7.069	22'-0"	264"	380,133
3'-6"	42"	9,621	22'-6"	270"	397.608
4'-0''	48"	12.566	23'-0"	276"	415.476
4'-6''	54"	15.904	23'-6''	282"	433.736
5'-0''	60''	19.635	24'-0''	288"	452.389
5'-6''	66"	23.758	24'-6''	294"	471.435
6'-0''	72"	28.274	25'-0"	300''	490.874
6'-6''	78"	33.183	25'-6"	306''	510.705
7'-0''	84"	38.485	26'-0''	312"	530.929
7'-6''	90"	44.179	26'-6"	318"	551.546
8'-0''	96"	50.266	27'-0''	324"	572.555
86.	102"	56.745	27'-6''	330"	593.957
9'-0''	108″	63.617	28'-0''	336"	615.752
9'-6''	114″	70.882	28'-6''	342"	637.940
10'-0''	120"	78.540	29'-0''	348"	660.520
10'-6''	126"	86.590	29'-6''	354"	683.493
11'-0"	132"	95.033	30'-0''	360''	706.858
11'-6"	138''	103.869	30'-6''	366"	730.617
12'-0'' 12'-6''	144" 150"	113.098	31′-0″	372"	754.768
	150	122.719	31'-6''	378"	779.311
13'-0"	156"	132.733	32'-0''	384"	804.248
13'-6''	162"	143.139	32'-6''	390"	829.577
14'-0''	168''	153.938	33'-0''	396″	855.299
14'-6"	174''	165.130	33'-6"	402''	881.413
15'-0"	180"	176.715	34'-0''	408''	907.920
15'-6''	186″	188.692	34'-6''	414"	934.820
16'-0''	192"	201.062	35'-0"	420''	962.113
16'-6''	198"	213.825	35'-6"	426"	989.798
17'-0''	204''	226.980	36'-0''	432''	1017.876
17'-6''	210"	240.528	36'-6''	438"	1046.347
18'-0''	216"	254.469	37'-0''	444"	1075.210
18'-6"	222"	268.803	37'-6"	450''	1104.46
19'-0''	228"	283.529	38'-0''	456"	1134.11
19'-6"	234″	298.648	38'-6"	462"	1164.15
20'-0''	240''	314.159	39'-0''	468"	1194.59
20'-6''	246"	330.064	39'-6"	476"	1225.41

Table 7-5Tower Diameter Versus Cross Sectional Areas

Table 7-6 Downcomer Dimensions



TABLES OF DOWNCOMER DIMENSIONS

| h/dia l/dia a _d /a _t |
|---|---|---|---|---|
| 0000. 0000. 0000. | .0200 .2800 .0048
.0205 .2834 .0050
.0210 .2868 .0051
.0215 .2901 .0053 | .0400 .3919 .0134 | .0600 .4750 .0245 | .0800 .5426 .0375 |
| .0005 .0447 .0000 | 0205 2834 0050 | .0400 .3919 .0134
.0405 .3943 .0137
.0410 .3966 .0139
.0415 .3989 .0142 | .0600 .4750 .0245
.0605 .4768 .0248
.0610 .4787 .0251
.0615 .4805 .0254
.0620 .4823 .0257 | .0805 .5441 .0378 |
| .0010 .0632 .0001 | 0210 2868 0051 | 0410 3966 0139 | .0610 .4787 .0251 | 0810 5457 0382 |
| .0015 .0774 .0001 | 0215 2901 0053 | 0415 3989 0142 | 0615 4805 0254 | 0815 5472 0385 |
| .0020 .0894 .0002 | .0220 .2934 .0055 | .0420 .4012 .0144 | 0620 4823 0257 | 0820 5487 0389 |
| .0020 .0894 .0002 | .0220 .2934 .0055 | | | |
| .0025 .0999 .0002
.0030 .1094 .0003
.0035 .1181 .0004
.0040 .1262 .0004
.0045 .1339 .0005 | .0225 .2966 .0057 | .0425 .4035 .0147
.0430 .4057 .0149
.0435 .4080 .0152
.0440 .4102 .0155
.0445 .4124 .0157 | .0625 .4841 .0260
.0630 .4859 .0263
.0635 .4877 .0266
.0640 .4895 .0270
.0645 .4913 .0273 | .0825 .5502 .0392 |
| 0020 1094 0003 | .0230 .2998 .0059 | 0430 4057 0149 | 0630 4859 0263 | .0830 .5518 .0396 |
| 0035 1181 0004 | .0235 .3030 .0061 | 0435 4080 0152 | .0635 .4877 .0266 | .0835 .5533 .0399 |
| 0040 1262 0004 | .0240 .3061 .0063 | 0440 4102 0155 | 0640 4895 0270 | 0840 5548 0403 |
| .0045 .1339 .0005 | .0245 .3092 .0065 | 0445 4124 0157 | 0645 4913 0273 | 0845 5563 0406 |
| .0045 .1559 .0005 | .0243 .3032 .0005 | | | |
| .0050 .1411 .0006 | .0250 .3122 .0067 | .0450 .4146 .0160 | .0650 .4931 .0276
.0655 .4948 .0279
.0660 .4966 .0282
.0665 .4983 .0285
.0670 .5000 .0288 | .0850 .5578 .0410 |
| | .0255 .3153 .0069 | 0455 4168 0162 | .0655 .4948 .0279 | .0855 .5592 .0413 |
| .0060 .1545 .0008 | 0260 3183 0071 | 0460 4190 0165 | .0660 .4966 .0282 | .0860 .5607 .0417 |
| .0065 .1607 .0009 | 0265 .3212 .0073 | .0465 .4211 .0168 | .0665 .4983 .0285 | .0865 .5622 .0421 |
| .0070 .1667 .0010 | .0255 .3153 .0069
.0260 .3183 .0071
.0265 .3212 .0073
.0270 .3242 .0075 | .0455 .4168 .0162
.0460 .4190 .0165
.0465 .4211 .0168
.0470 .4233 .0171 | .0670 .5000 .0288 | .0870 .5637 .0424 |
| | | | | |
| .0075 .1726 .0011 | .0275 .3271 .0077 | .0475 .4254 .0173 | .0675 .5018 .0292 | .0875 .5651 .0428 |
| .0080 .1782 .0012 | .0280 .3299 .0079 | .0480 .4275 .0176 | .0680 .5035 .0295 | .0880 .5666 .0431 |
| .0085 .1836 .0013 | .0285 .3328 .0081 | .0485 .4296 .0179 | .0685 .5052 .0298 | .0885 .5680 .0435 |
| .0090 .1889 .0014 | .0290 .3356 .0083 | .0490 .4317 .0181 | .0690 .5069 .0301 | .0890 .5695 .0439 |
| .0095 .1940 .0016 | .0275 .3271 .0077
.0280 .3299 .0079
.0285 .3328 .0081
.0290 .3356 .0083
.0295 .3384 .0085 | .0495 .4338 .0184 | .0675 .5018 .0292
.0680 .5035 .0295
.0685 .5052 .0298
.0690 .5069 .0301
.0695 .5086 .0304 | .0895 .5709 .0442 |
| | | | | |
| 0100 1000 0017 | .0300 .3412 .0087
.0305 .3439 .0090
.0310 .3466 .0092
.0315 .3493 .0094
.0320 .3520 .0096 | 0500 4350 0187 | 0700 5102 0209 | 0000 5724 0446 |
| .0100 .1990 .0017
.0105 .2039 .0018 | .0300 .3412 .0087 | .0500 .4359 .0187 | 0705 5103 0308 | 0005 5724 0440 |
| .0110 .2086 .0020 | .0305 .3439 .0090 | .0505 .4379 .0190 | 0710 5126 0314 | .0905 .5758 .0449 |
| .0115 .2132 .0021 | .0310 .3466 .0092 | .0510 .4400 .0193 | 0715 5150 0014 | .0910 .5752 .0453 |
| .0120 .2178 .0022 | 0315 .3493 .0094 | .0515 .4420 .0195 | .0700 .5103 .0308
.0705 .5120 .0311
.0710 .5136 .0314
.0715 .5153 .0318
.0720 .5170 .0321 | .0915 .5788 .0457 |
| .0120 .2178 .0022 | .0320 .3320 .0098 | .0320 .4441 .0198 | .0720 .5170 .0321 | .0920 .3781 .0400 |
| .0125 .2222 .0024 | .0325 .3546 .0098
.0330 .3573 .0101
.0335 .3599 .0103
.0340 .3625 .0105 | .0525 .4461 .0201 | .0725 .5186 .0324
.0730 .5203 .0327
.0735 .5219 .0331
.0740 .5235 .0334
.0745 .5252 .0337 | .0925 .5795 .0464 |
| .0130 .2265 .0025 | .0330 .3573 .0101 | .0530 .4481 .0204 | .0730 .5203 .0327 | .0930 .5809 .0468 |
| .0135 .2308 .0027 | .0335 .3599 .0103 | .0535 .4501 .0207 | .0735 .5219 .0331 | .0935 .5823 .0472 |
| .0140 .2350 .0028 | .0340 .3625 .0105 | .0540 .4520 .0210 | .0740 .5235 .0334 | .0940 .5837 .0475 |
| .0145 .2391 .0030 | .0345 .3650 .0108 | .0545 .4540 .0212 | .0745 .5252 .0337 | .0945 .5850 .0479 |
| | | | | |
| .0150 .2431 .0031 | .0350 .3676 .0110 | .0550 .4560 .0215 | .0750 .5268 .0341
.0755 .5284 .0344
.0760 .5300 .0347
.0765 .5316 .0351
.0770 .5332 .0354 | .0950 .5864 .0483
.0955 .5878 .0486
.0960 .5892 .0490
.0965 .5906 .0494
.0970 .5919 .0498 |
| .0155 .2471 .0033 | .0355 .3701 .0112 | .0555 .4579 .0218 | .0755 .5284 .0344 | .0955 .5878 .0486 |
| .0160 .2510 .0034 | .0360 .3726 .0115 | .0560 .4598 .0221 | .0760 .5300 .0347 | .0960 .5892 .0490 |
| .0165 .2548 .0036 | .0365 .3751 .0117 | .0565 .4618 .0224 | .0765 .5316 .0351 | .0965 .5906 .0494 |
| .0170 .2585 .0037 | .0350 .3676 .0110
.0355 .3701 .0112
.0360 .3726 .0115
.0365 .3751 .0117
.0370 .3775 .0119 | .0570 .4637 .0227 | .0770 .5332 .0354 | .0970 .5919 .0498 |
| 0175 0000 0000 | 0005 0000 0100 | 0575 4656 0000 | 0775 5340 0258 | 0075 5022 0501 |
| | .0375 .3800 .0122 | .0575 .4656 .0230
.0580 .4675 .0233
.0585 .4694 .0236
.0590 .4712 .0230
.0595 .4731 .0242 | .0775 .5348 .0358 | .0975 .5933 .0501
.0980 .5946 .0505
.0985 .5960 .0509
.0990 .5973 .0513 |
| .0180 .2659 .0041 | .0380 .3824 .0124 | .0580 .46/5 .0233 | .0780 .5363 .0361 | .0980 .5946 .0505 |
| .0185 .2695 .0042 | .0385 .3848 .0127 | .0585 .4694 .0236 | .0785 .5379 .0364 | .0985 .5960 .0509 |
| .0190 .2730 .0044 | .0390 .3872 .0129 | .0590 .4/12 .0230 | .0790 .5395 .0368 | .0990 .5973 .0513 |
| .0195 .2765 .0046 | .0395 .3896 .0132 | .0595 .4731 .0242 | .0795 .5410 .0371 | .0995 .5987 .0517 |
| | | | | |

Table 7-7Downcomer Dimensions

H/DIA L/DIA A _d /A _t				φ.,
.1000 .6000 .0520	1275 6671 0743	1550 7238 0986	.1825 .7725 .1249 .1830 .7733 .1253 .1835 .7742 .1258 .1840 .7750 .1263 .1845 .7758 .1268	2100 8146 1527
.1000 .6000 .0520	1200 6692 0743	1550 .7258 .0980	1920 7722 1249	.2100 .8146 .152/
.1005 .6013 .0524	.1280 .0002 .0747	.1555 .7248 .0991	.1830 .7733 .1253	.2105 .8153 .1532
.1010 .6027 .0528	.1285 .6693 .0751	.1560 ./25/ .0996	.1835 .//42 .1258	.2110 .8160 .1537
.1015 .6040 .0532	.1290 .6704 .0755	.1565 .7267 .1000	.1840 .7750 .1263	.2115 .8167 .1542
	.1275 .6671 .0743 .1280 .6682 .0747 .1285 .6693 .0751 .1290 .6704 .0755 .1295 .6715 .0760			
.1025 .6066 .0540	.1300 .6726 .0764 .1305 .6737 .0768 .1310 .6748 .0773 .1315 .6759 .0777 .1320 .6770 .0781	.1575 .7285 .1009	.1850 .7766 .1273	.2125 .8182 .1553
.1030 .6079 .0544	.1305 .6737 .0768	.1580 .7295 .1014	.1855 .7774 .1278	.2130 .8189 .1558
1035 .6092 .0547	.1310 .6748 .0773	.1585 .7304 .1019	.1860 .7782 .1283	2135 8196 1563
1040 .6105 .0551	.1315 .6759 .0777	1590 7314 1023	1865 7790 1288	2140 8203 1568
.1045 .6118 .0555	.1320 .6770 .0781	.1595 .7323 .1028	.1870 .7798 .1293	.2145 .8210 .1573
.1050 .6131 .0559	.1325 .6781 .0785 .1330 .6791 .0790 .1335 .6802 .0794 .1340 .6813 .0798 .1345 .6824 .0803	.1600 .7332 .1033	.1875 .7806 .1298 .1880 .7814 .1303 .1885 .7822 .1308 .1890 .7830 .1313 .1895 .7838 .1318	2150 8216 1579
.1055 .6144 .0563	1330 6791 0790	1605 7341 1037	1880 7814 1303	.2150 .8216 .1579 .2155 .8223 .1584
1060 6157 0567	1335 6802 0794	1610 7351 1042	1995 7922 1309	.2160 .8230 .1589
.1060 .6157 .0567 .1065 .6170 .0571	1340 6813 0708	1616 7360 1047	1900 7920 1212	.2165 .8237 .1594
.1070 .6182 .0575	1345 6934 0003	1000 7300 1047	1090 .7030 .1313	.2170 .8244 .1600
.1075 .6195 .0579	.1350 .6834 .0807 .1355 .6845 .0811 .1360 .6856 .0816 .1365 .6866 .0820 .1370 .6877 .0825	.1625 .7378 .1056	.1900 .7846 .1323	.2175 .8251 .1605
.1080 .6208 .0583	.1355 .6845 .0811	.1630 .7387 .1061	.1905 .7854 .1328	.2180 .8258 .1610
.1085 .6220 .0587	.1360 .6856 .0816	.1635 .7396 .1066	.1910 .7862 .1333	.2185 .8265 .1615
.1090 .6233 .0591	.1365 .6866 .0820	.1640 .7406 .1070	.1915 .7870 .1338	.2190 .8271 .1621
1095 6245 0595	1370 6877 0825	1645 7415 1075	1920 7877 1343	2195 8278 1626
,				
.1100 .6258 .0598	1275 6997 0920	1650 7424 1080	1075 7885 1249	.2200 .8285 .1631 .2205 .8292 .1636 .2210 .8298 .1642 .2215 .8305 .1647
.1105 .6270 .0602	.1375 .6887 .0829 .1380 .6898 .0833 .1385 .6908 .0838 .1390 .6919 .0842 .1395 .6929 .0847	1655 7424 1080	.1925 .7885 .1348 .1930 .7893 .1353 .1935 .7901 .1358 .1940 .7909 .1363 .1945 .7916 .1368	2200 .0205 .1031
.1105 .6270 .0602	.1380 .0898 .0833	1000 7433 1084	1025 7001 1259	.2205 .8292 .1636
.1110 .6283 .0606	.1365 .0010 .0010	.1660 .7442 .1089	.1935 ./901 .1358	.2210 .8298 .1642
.1115 .6295 .0610	.1390 .6919 .0842	.1665 .7451 .1094	.1940 .7909 .1363	.2215 .8305 .164/
				.2220 .8312 .1652
.1125 .6320 .0619	.1400 .6940 .0851	.1675 .7468 .1103	.1950 .7924 .1373	.2225 .8319 .1658
.1130 .6332 .0623	.1405 .6950 .0855	.1680 .7477 .1108	.1955 .7932 .1378	.2230 .8325 .1663
.1135 .6344 .0627	.1410 .6960 .0860	.1685 .7486 .1113	.1960 .7939 .1383	.2235 .8332 .1668
.1140 .6356 .0631	.1415 .6971 .0864	.1690 .7495 .1118	.1965 .7947 .1388	.2240 .8338 .1674
.1145 .6368 .0635	.1400 .6940 .0851 .1405 .6950 .0855 .1410 .6960 .0860 .1415 .6971 .0864 .1420 .6981 .0869	.1695 .7504 .1122	.1970 .7955 .1393	.2245 .8345 .1679
.1150 .6380 .0639	.1425 .6991 .0873	.1700 .7513 .1127	.1975 .7962 .1398	.2250 .8352 .1684 .2255 .8358 .1689 .2260 .8365 .1695
.1155 .6392 .0643	,1430 ,7001 ,0878	.1705 .7521 .1132	.1980 .7970 .1403	.2255 .8358 .1689
.1160 .6404 .0647	.1435 .7012 .0882	.1710 .7530 .1137	.1985 .7977 .1409	2260 8365 1695
.1165 .6416 .0651	.1440 .7022 .0886	.1715 .7539 .1142	.1990 .7985 .1414	.2265 .8371 .1700
.1170 .6428 .0655	.1425 .6991 .0873 .1430 .7001 .0878 .1435 .7012 .0882 .1440 .7022 .0886 .1445 .7032 .0891	.1720 .7548 .1146	.1995 .7992 .1419	.2270 .8378 .1705
	.1450 .7042 .0895 .1455 .7052 .0900 .1460 .7062 .0904 .1465 .7072 .0909 .1470 .7082 .0913	.1725 .7556 .1151 .1730 .7565 .1156 .1735 .7574 .1161 .1740 .7582 .1166 .1745 .7591 .1171	.2000 .8000 1424	.2275 .8384 .1711 .2280 .8391 .1716 .2285 .8397 .1721 .2290 .8404 .1727
.1180 .6452 .0663	1455 7052 0900	1730 7565 1156	2005 8007 1429	2280 8301 1716
.1185 .6464 .0667	1460 7062 0904	1735 7574 1161	2010 8015 1424	2285 8207 1721
.1190 .6476 .0671	1465 7072 0000	1740 7582 1166	2015 8022 1429	2203 .0357 .1721
.1195 .6488 .0676	1470 7082 0913	1745 7501 1171	2020 8030 1444	.2295 .8410 .1732
.1200 .6499 .0680	.1475 .7092 .0918	.1750 .7599 .1175	.2025 .8037 .1449	.2300 .8417 .1738
.1205 .6511 .0684	.1480 .7102 .0922	.1755 .7608 .1180	.2030 .8045 .1454	.2305 .8423 .1743
.1210 .6523 .0688	.1485 .7112 .0927	.1760 .7616 .1185	.2035 .8052 .1460	.2310 .8429 .1748
.1215 .6534 .0692	.1490 .7122 .0932	.1765 .7625 .1190	.2040 .8059 .1465	.2315 .8436 .1754
.1220 .6546 .0696	.1475 .7092 .0918 .1480 .7102 .0922 .1485 .7112 .0927 .1490 .7122 .0932 .1495 .7132 .0936	.1770 .7633 .1195	.2045 .8067 .1470	.2320 .8442 .1759
.1230 .6569 .0705	1505 7151 0945	.1780 .7650 1204	.2055 .8081 .1480	2330 8455 1770
.1235 .6580 .0709	1510 7161 0950	1785 7659 1209	.2060 .8089 .1485	2335 8461 1775
.1240 .6592 .0713	1515 7171 0954	1790 7667 1214	2065 8096 1490	2340 8467 1791
.1245 .6603 .0717	.1500 .7141 .0941 .1505 .7151 .0945 .1510 .7161 .0950 .1515 .7171 .0954 .1520 .7180 .0959	.1775 .7642 .1200 .1780 .7650 .1204 .1785 .7659 .1209 .1790 .7667 .1214 .1795 .7675 .1219	.2065 .8096 .1490 .2070 .8103 .1496	.2325 .8449 .1764 .2330 .8455 .1770 .2335 .8461 .1775 .2340 .8467 .1781 .2345 .8474 .1786
.1250 .6614 .0721	.1525 .7190 .9063 .1530 .7200 .0968 .1535 .7209 .0973 .1540 .7219 .0977 .1545 .7229 .0982	1000 ./084 .1224	.2075 .8110 .1501 .2080 .8118 .1506 .2085 .8125 .1511 .2090 .8132 .1516 .2095 .8139 .1521	.2350 .8480 .1791
.1255 .6626 .0726	.1530 .7200 .0968	.1805 ./692 .1229	.2080 .8118 .1506	.2355 .8486 .1797
.1260 .6637 .0730	.1535 ./209 .0973	.1810 .7700 .1234	.2085 .8125 .1511	.2360 .8492 .1802
.1265 .6648 .0734	.1540 .7219 .0977	.1815 .7709 .1239	.2090 .8132 .1516 .2095 .8139 .1521	.2365 .8499 .1808
.1270 .6659 .0738	.1545 .7229 .0982	.1820 .7717 .1244	.2095 .8139 .1521	.2370 .8505 .1813

Table 7-8Downcomer Dimensions

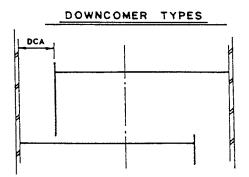
H/DIA L/DIA A _d /A _t	H/DIA L/DIA A _d /A _t	H/DIA L/DIA A _d /A _t	H/DIA L/DIA A _d /A _t	H/DIA L/DIA A _d /A _t
0075 0511 1010	2650 8827 2122	.2925 .9098 .2436 .2930 .9103 .2442 .2935 .9107 .2448 .2940 .9112 .2453 .2945 .9116 .2459	3200 9330 2759	3475 9524 2080
.23/5 .0511 .1010	2655 8832 2128	2930 9103 2442	3205 9333 2765	3490 9527 3005
2380 .8517 .1624	2560 8837 2133	2935 9107 2448	3210 9337 2771	3485 9530 3101
2385 .8523 .1829	200 .003/ .2133	2040 0112 2452	2216 2341 2777	3485 .5550 .3101
2390 .8529 .1835	2005 .0045 .2135	2045 0116 2450	3213 .3341 .2777	.3490 .9533 .3107
.2400 .8542 .1845	.2675 .8853 .2150 .2680 .8858 .2156 .2685 .8864 .2161 .2690 .8869 .2167 .2695 .8874 .2173	.2950 .9121 .2465	.3225 .9349 .2788 .3230 .9352 .2794 .3235 .9356 .2800 .3240 .9360 .2806 .3245 .9364 .2812	.3500 .9539 .3119
.2405 .8548 .1851	.2680 .8858 .2156	.2955 .9125 .2471	.3230 .9352 .2794	.3505 .9543 .3125
,2410 .8554 .1856	.2685 .8864 .2161	.2960 .9130 .2477	.3235 .9356 .2800	.3510 .9546 .3131
.2415 .8560 .1862	.2690 .8869 .2167	.2965 .9134 .2482	.3240 .9360 .2806	.3515 .9549 .3137
.2420 .8566 .1867	.2695 .8874 .2173			
.2425 .8572 .1873	.2700 .8879 .2178	.2975 .9143 .2494	.3250 .9367 .2818 .3255 .9371 .2824 .3260 .9375 .2830 .3265 .9379 .2511 .3270 .9382 .2842	.3525 .9555 .3150
.2430 .8578 .1878	.2705 .8884 .2184	.2980 .9148 .2500	.3255 .9371 .2824	.3530 .9558 .3156
.2435 .8584 .1884	.2710 .8890 .2190	.2985 .9152 .2506	.3260 .9375 .2830	.3535 .9561 .3162
2440 8590 1889	.2715 .8895 .2195	2990 9156 2511	3265 9379 2511	3540 9564 3168
.1445 .8596 .1895	.2700 .8879 .2178 .2705 .8884 .2184 .2710 .8890 .2190 .2715 .8895 .2195 .2720 .8900 .2201	.2995 .9161 .2517	.3270 .9382 .2842	.3545 .9567 .3174
.2450 .8602 .1900 .2455 .8608 .1906	2730 8905 2207	3000 9165 2523	.3275 .9386 .2848	3555 9573 3180
.2460 .8614 .1911	2735 8015 2218	2010 9174 2535	3285 9393 2860	3560 9576 2192
2466 8610 1017	2740 8020 22210	2015 0179 2541	3200 9397 2866	3565 9570 3192
.2465 .8619 .1917 .2470 .8625 .1922	.2745 .8925 .2229	.3000 .9165 .2523 .3005 .9170 .2529 .3010 .9174 .2535 .3015 .9178 .2541 .3020 .9138 .2547	.3295 .9401 .2872	.3570 .9582 .3204
.2475 .8631 .1927	.2750 .8930 .2235	.3025 .9187 .2552	.3300 .9404 .2878	.3575 .9585 .3211
.2480 .8637 .1933	.2755 .8935 .2241	.3030 .9191 .2558	.3305 .9408 .2884	.3580 .9588 .3217
.2485 .8643 .1938	.2760 .8940 .2246	.3035 .9195 .2564	.3310 .9411 .2890	.3585 .9591 .3223
.2490 .8649 .1944	.2765 .8945 .2252	.3040 .9200 .2570	.3315 .9415 .2896	.3590 .9594 .3229
.2495 .8654 .1949	.2770 .8950 .2258	.3025 .9187 .2552 .3030 .9191 .2558 .3035 .9195 .2564 .3040 .9200 .2570 .3045 .9204 .2576	.3320 .9419 .2902	.3595 .9597 .3235
.2500 .8660 .1955	.2775 .8955 .2264	.3050 .9208 .2582 .3055 .9212 .2588 .3060 .9217 .2593 .3065 .9221 .2599 .3070 .9225 .2605	.3325 .9422 .2908	.3600 .9600 .3241
.2505 .8666 .1961	2780 .8960 .2269	3055 .9212 .2588	.3330 .9426 .2914	.3605 .9603 .3247
.2510 .8672 .1966	2785 8965 2275	.3060 .9217 .2593	.3335 .9429 .2920	3610 9606 3253
.2515 .8678 .1972	2790 .8970 .2281	3065 9221 2599	.3340 .9433 .2926	3615 9609 3259
.2515 .8678 .1972 .2520 .8683 .1977	.2795 .8975 .2286	.3070 .9225 .2605	.3345 .9436 .2932	.3620 .9612 .3265
.2530 .8695 .1988	2005 2005 2202	.3075 .9229 .2617	2225 0442 2044	.3025 .9014 .3272
.2535 .8700 .1994	.2805 .8985 .2298	.3080 .9233 .2017	.3335 .9443 .2944	.3630 .9617 .3278
.2540 .8706 .1999	2810 .8990 .2304	3000 0242 2623	2265 0450 2056	.3035 .9020 .3284
.2545 .8712 .2005	.2820 .8999 .2315	.3075 .9229 .2611 .3080 .9233 .2617 .3085 .9237 .2623 .3090 .9242 .2629 .3095 .9246 .2635	.3370 .9454 .2962	.3645 .9626 .3296
.2550 .8717 .2010 .2555 .8723 .2016	.2825 .9004 .2321	.3100 .9250 .2640 .3105 .9254 .2646 .3110 .9258 .2652 .3115 .9262 .2658	.3375 .9457 .2968	.3650 .9629 .3302
.2555 .8723 .2016	.2830 .9009 .2326	.3105 .9254 .2646	.3380 .9461 .2974	.3655 .9631 .3308
.2560 .8728 .2021	.2835 .9014 .2332	.3110 .9258 .2652	.3385 .9464 .2980	.3660 .9634 .3315
				.3665 .9637 .3321
.2575 .8745 .2038	.2850 .9028 .2349	.3125 .9270 .2670 .3130 .9274 .2676 .3135 .9278 .2682 .3140 .9282 .2688 .3145 .9286 .2693	.3400 .9474 .2998	.3675 .9642 .3333
.2575 .8745 .2038 .2580 .8751 .2044	.2855 .9033 .2355	.3130 .9274 .2676	.3405 .9478 .3004	3680 9645 3339
.2585 .8756 .2049	.2860 .9038 .2361	.3135 .9278 .2682	.3410 .9481 .3010	3685 9648 3345
.2590 .8762 .2055	2865 9043 2367	3140 9282 2688	3415 9484 3016	3690 9651 3351
.2595 .8767 .2060	.2870 .9047 .2872	.3145 .9286 .2693	.3420 .9488 .3022	.3675 .9642 .3333 .3680 .9645 .3339 .3685 .9648 .3345 .3690 .9651 .3351 .3695 .9653 .3357
.2000 .8//3 .2066	.28/5 .9052 .2378 2880 9057 2384	.3150 .9290 .2699	.3425 .9491 .3028	.3/00 .9656 .3364
2610 8784 2077	2885 0061 2200	3160 9298 2711	3435 9494 3034	3710 0661 3370
2615 8789 2083	2800 0066 2205	3165 9302 2717	3440 9501 3046	3715 0664 220
.2620 .8794 .2088	.2895 .9071 .2401	.3150 .9290 .2699 .3155 .9294 .2705 .3160 .9298 .2711 .3165 .9302 .2717 .3170 .9306 .2723	.3445 .9504 .3053	.3700 .9656 .3364 .3705 .9659 .3370 .3710 .9661 .3374 .3715 .9664 .338 .3720 .9667 .3384
.2625 .8800 .2094	.2900 .9075 .2407	.3175 .9310 .2729 .3180 .9314 .2735 .3185 .9318 .2741 .3190 .9322 .2747 .3195 .9326 .2753	.3450 .9507 .3059	.3750 .9669 .339
.2030 .8805 .2100	.2905 .9080 .2413	,3180 ,9314 ,2735	.3455 .9511 .3065	.3/30 .9672 .340
.2635 .8811 .2105	.2910 .9084 .2419	.3185 .9318 .2741	.3460 .9514 .3071	.3735 .9676 .340
.2640 .8816 .2111	.2915 .9089 .2424	.3190 .9322 .2747	.3465 .9517 .3077	.3740 .9677 .341

Table 7-9 Downcomer Dimensions

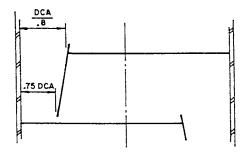
H/DIA L/DIA A _d /A _t	H/DIA L/DIA A _d /A _t	H/DłA L/DIA A _d /A _t	H/DIA L/DIA A _d /A _t	H/DIA L/DIA A _d /A _t
.3750 .9682 .3425	4000 9798 3735	4250 9887 4049	4500 9950 4364	4750 9987 4682
.3755 .9685 .3431	4005 9800 3742	4255 9888 4055	4505 .9951 .4371	4755 .9988 .4688
.3760 .9688 .3438	4010 .9802 .3748	.4260 .9890 .4061	.4510 .9952 .4377	.4760 .9988 .4695
.3765 .9690 .3444	.4015 .9804 .3754	.4265 .9891 .4068	.4515 .9953 .4383	.4765 .9989 .4701
.3770 .9693 .3450	.4020 .9806 .3760	.4270 .9893 .4074	.4500 .9950 .4364 .4505 .9951 .4371 .4510 .9952 .4377 .4515 .9953 .4383 .4520 .9954 .4390	.4770 .9989 .4707
.3775 .9695 .3456	4025 9808 3767	4275 9894 4080	4525 9955 4396	4775 9990 4714
3780 9698 3462	4030 .9810 .3773	4280 .9896 .4086	4530 .9956 .4402	4780 9990 4720
.3780 .9698 .3462 .3785 .9700 .3468 .3790 .9703 .3475	.4035 .9812 .3779	.4285 .9897 .4093	.4535 .9957 .4409	.4785 .9991 .4726
3790 9703 3475	.4040 .9814 .3785	4290 9899 4099	4540 .9958 .4415	4790 9991 4733
.3795 .9705 .3481	.4045 .9816 .3791	.4295 .9900 .4105	.4525 .9955 .4396 .4530 .9956 .4402 .4535 .9957 .4409 .4540 .9958 .4415 .4545 .9959 .4421	.4795 .9992 .4739
.3800 .9708 .3487	4050 0010 0700	4000 0000 4110	.4550 .9959 .4428 .4555 .9960 .4434 .4560 .9961 .4440 .4565 .9962 .4447 .4570 .9963 .4453	4000 0000 4745
2805 0710 2402	4050 .9810 .3798	4300 .9902 .4112	4550 .9959 .4428	.4800 .9992 .4745
.3805 .9710 .3493 .3810 .9713 .3499 .3815 .9715 .3505	4060 0822 3810	4310 0004 4124	4555 .9960 .4434	4810 0003 4758
3815 9715 3505	4065 9824 3816	4215 9906 4131	4565 0062 4440	4815 0003 4765
.3820 .9718 .3512	.4070 .9825 .3823	.4320 .9907 .4137	.4570 .9963 .4453	.4820 .9994 .4771
.3825 .9720 .3518	.4075 .9827 .3829	.4325 .9908 .4143	.4575 .9964 .4460 .4580 .9965 .4466 .4585 .9965 .4472 .4590 .9966 .4479 .4595 .9967 .4485	.4825 .9994 .4777
.3830 .9722 .3524 .3835 .9725 .3530	.4080 .9829 .3835	.4330 .9910 .4149	.4580 .9965 .4466	.4830 .9994 .4784
.3835 .9725 .3530	.4085 .9831 .3842	.4335 .9911 .4156	.4585 .9965 .4472	.4835 .9995 .4790
.3840 .9727 .3536	.4090 .9833 .3848	.4340 .9912 .4162	.4590 .9966 .4479	.4840 .9995 .4796
.3850 .9732 .3549	.4100 .9837 .3860	.4350 .9915 .4175	.4600 .9968 .4491	.4850 .9995 .4809
.3855 .9734 .3555	.4105 .9838 .3867	.4355 .9916 .4181	.4605 .9969 .4498	.4855 .9996 .4815
.3860 .9737 .3561	.4110 .9840 .3873	.4360 .9918 .4187	.4610 .9970 .4505	.4860 .9996 .4822
.3865 .9739 .3567	.4115 .9842 .3879	.4365 .9919 .4194	.4615 .9970 .4510	.4865 .9996 .4828
.3870 .9741 .3574	.4120 .9844 .3885	.4370 .9920 .4200	.4600 .9968 .4491 .4605 .9969 .4498 .4610 .9970 .4505 .4615 .9970 .4510 .4620 .9971 .4517	.4870 .9997 .4834
.3880 .9746 .3586 .3885 .9748 .3592	.4130 .9847 .3898	.4380 .9923 .4213	.4630 .9973 .4529	4880 .9997 .4847
.3885 .9748 .3592	.4135 .9849 .3904	.4385 .9924 .4219	.4635 .9973 .4536	.4885 .9997 .4854
.3890 .9750 .3598	.4140 .9851 .3910	.4390 .9925 .4225	.4640 .9974 .4542	.4890 .9998 .4860
.3895 .9753 .3605	.4145 .9853 .3917	.4395 .9927 .4232	.4625 .9972 .4523 .4630 .9973 .4529 .4635 .9973 .4536 .4640 .9974 .4542 .4645 .9975 .4548	.4895 .9998 .4866
.3900 .9755 .3611	.4150 .9854 .3923	.4400 .9928 .4238	.4650 .9975 .4555	.4900 .9998 .4873
.3905 .9757 .3617	.4155 .9856 .3929	.4405 .9929 .4244	.4655 .9976 .4561	.4905 .9998 .4879
.3910 .9759 .3623	.4160 .9858 .3936	.4410 .9930 .4251	.4660 .9977 .4567	.4910 .9998 .4885
.3915 .9762 .3629	.4165 .9860 .3942	.4415 .9931 .4257	.4665 .9978 .4574	.4915 .9999 .4892
.3920 .9764 .3636	.4170 .9861 .3948	.4420 .9932 .4263	.4650 .9975 .4555 .4655 .9976 .4561 .4660 .9977 .4567 .4665 .9978 .4574 .4670 .9978 .4580	.4920 .9999 .4898
.3925 .9766 .3642	.4175 .9863 .3642	.4425 .9934 .4270	.4675 .9979 .4568 .4680 .9979 .4593 .4685 .9980 .4599 .4690 .9981 .4606 .4695 .9981 .4612	.4925 .9999 .4905
.3930 .9768 .3648	.4180 .9865 .3961	.4430 .9935 .4276	.4680 .9979 .4593	.4930 .9999 .4911
.3935 .9771 .3654	.4185 .9866 .3967	.4435 .9936 .4282	.4685 .9980 .4599	.4935 .9999 .4917
.3940 .9773 .3661	.4190 .9868 .3973	.4440 .9937 .4288	.4690 .9981 .4606	.4940 .9999 .4924
.3945 .9775 .3667	.4195 .9870 .3979	.4445 .9938 .4295	.4695 .9981 .4612	.4945 .9999 .4930
.3950 .9777 .3673	.4200 .9871 .3986	.4450 .9939 .4301	.4700 .9982 .4618	.4950 1.0000 .4936
.3955 .9779 .3679	.4205 .9873 .3992	.4455 .9940 .4307	.4705 .9983 .4625	.4955 1.0000 .4943
.3960 .9781 .3685	.4210 .9874 .3998	.4460 .9942 .4314	.4710 .9983 .4631	.4960 1.0000 .4949
.3965 .9783 .3692	.4215 .9876 .4005	.4465 .9943 .4320	.4715 .9984 .4637	.4965 1.0000 .4955
.3970 .9786 .3698	.4220 .9878 .4011	.4470 .9944 .4326	.4700 .9982 .4618 .4705 .9983 .4625 .4710 .9983 .4631 .4715 .9984 .4637 .4720 .9984 .4644	.4970 1.0000 .4962
3980 9790 3710	4230 9881 4023	.4480 .9946 .4339	.4725 .9985 .4650 .4730 .9985 .4656 .4735 .9986 .4663 .4740 .9986 .4669 .4745 .9987 .4675	4980 1.0000 .4975
3985 9792 3717	4235 9882 4030	.4485 .9947 .4345	.4735 .9986 .4663	.4985 1.0000 .4981
.3990 .9794 .3723	4240 9884 4036	.4490 .9948 .4352	.4740 .9986 .4669	.4990 1.0000 .4987
.3995 .9796 .3729	.4245 .9885 .4042	.4495 .9949 .4358	.4745 .9987 .4675	.4995 1.0000 .4994
	2.2			.5000 1.0000 .5000
	· · · · · · · · · · · · · · · · · · ·			

Table 7-10Determining Number of Passes

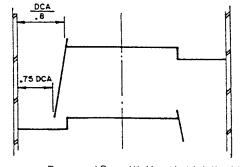
Tray Spacing (inch)	Increase No. of Passes gpm/in. of Weir Exceeds		
12	3		
18	18		



Vertical Downcomer — Flat Deck: Used as a standard unless process reasons dictate otherwise

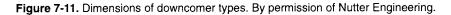


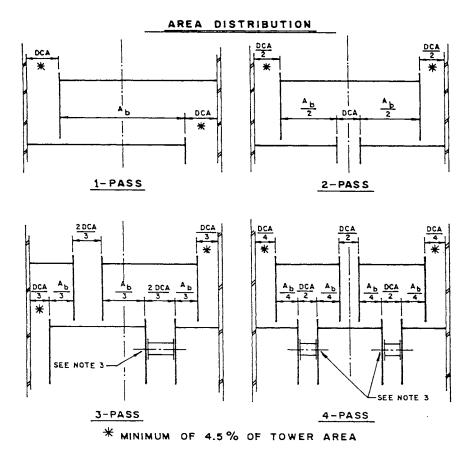
Sloped Downcomer — Flat Deck: Used in high pressure systems where vapor disengagement out of the top portion of the downcomer is critical.



Sloped Downcomer — Recessed Pan: (1) Used in high liquid rate systems to help alleviate hydraulic problems (2) Used in low liquid rate systems to

maintain positive downcomer seal





NOTES:

- Trays are designed using as few passes as possible. Increasing number of passes decreases efficiency and increases cost.
- (2) Multipass trays are designed with equal-bubbling areas. Weir lengths or heights can be adjusted to provide uniform tray hydraulics.
- (3) Vapor equalizers are a recommended option to guarantee vapor equalization between compartments.

Figure 7-12. Area distribution for different pass types. By permission of Nutter Engineering.

(text continued from page 492)

D = tower diameter, ftFoam fac. = foam factor (see Table 7-4) $Q_L = \text{volumetric liquid flow rate, ft}^3/\text{s}$ $Q_V = \text{volumetric vapor flow rate, ft}^3/\text{s}$ $R_D = \text{density radical}$ $S_{fac} = \text{safety factor}$ $T_{SF} = \text{tray spacing factor}$ V = velocity at zero liquid load, ft/sec $W_L = \text{liquid flow rate, lb/h}$ $W_V = \text{vapor flow rate, lb/h}$

Greek Letters

 ρ_L = liquid density, lb/ft³ ρ_V = vapor density, lb/ft³ σ = surface tension, dvnes/cm

PACKED TOWER DESIGN

Introduction

Packed towers are used increasingly in a variety of applications in the chemical process industries. The reasons for this are an increase in the availability of design information, the evolution of higher-capacity and higher-efficiency mass-transfer packings, and improvements in distributors and support plates. Today, packings can be considered for most services where high numbers of theoretical stages are required for mass transfer.

Most of the fractionating columns in gas-processing plants that were previously equipped with trays can now be filled with packings. Packings, such as slotted rings in metal and plastic and plastic saddles, offer large improvements in capacity for absorption and heat transfer applications. An example of this is the use of slotted rings in a gas-plant demethanizer, which resulted in a significant capacity improvement over trays.

Packed towers are used increasingly in small fractionating towers and absorbers where accessibility is restricted. Their use is also advantageous where corrosion control is essential. Because of liquid-distribution concerns, packing still is usually not recommended for large-diameter columns or where a high turndown ratio is required. All packings tend to spread liquid as it flows downwards. This characteristic is adequate in dispersing the liquid between distribution points, but not to correct poor distribution across a tower without considerable loss of efficiency. Studies of liquid spread in tower packing have been reported by Hoak [14]. Figure 7-13 illustrates the sections of a packed tower.

Pressure Drop

The pressure drop (ΔP) through a packed bed represents the frictional loss, kinetic energy loss through the packing, and the force exerted by the operating liquid holdup. At constant ΔP , the packed bed has less volumetric liquid holdup in high-liquid-density systems. But with lowdensity liquids, the volumetric liquid holdup can be significantly greater than for water at the same ΔP . At atmospheric pressure and in vacuum services, the static head produced by the gas is negligible. In highpressure fractionators requiring large number of theoretical stages, the static head of vapor can be considerable from the condenser to the reboiler. Here, the value should be added to the calculated ΔP in determining the bottom column pressure and reboiler temperature. Pressure drop may be estimated from Figure 7-14 for the particular packing system properties and operating conditions [15], or may be calculated by the following equation as represented by Leva [16].

$$\Delta \mathbf{P} = \alpha 10^{\beta L} \left(\frac{\mathbf{G}_{\mathrm{I}}^2}{\rho_{\mathrm{G}}} \right) \tag{7-61}$$

The equation was originally developed for air-water systems. For other liquids, however, L is multiplied by the ratio of the density of water to the density of liquid. Table 7-11 shows ΔP correlations for different types of packing material [17].

Flooding

In packed columns, flooding is caused when the energy potential of the vapor is greater than that of the liquid (that is, when all the void spaces in the packing are filled with liquid), so that the liquid travels up and the column floods. This gives a ΔP of 2–3 in. H₂O/ft of packing depth, and a column cannot be operated successfully under these conditions. Flooding depends on a number of parameters, and empirical correlations of the variables have been presented by Bain and Hougen [18], Lobo *et al.* [19], Sherwood *et al.* [20], and Eckert [21].

Eckert's modification of Sherwood's flooding correlation has been widely applied for designing packed columns. Blackwell [13] developed

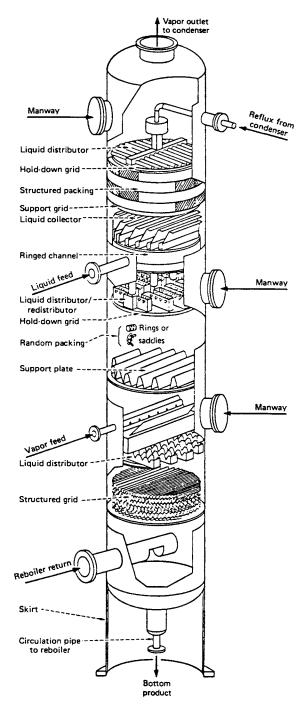


Figure 7-13. Packed tower and internals. By permission of Chemical Engineering.

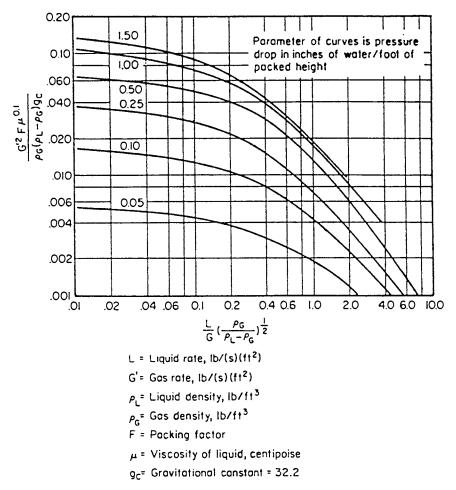


Figure 7-14. Correlation of flow rate and pressure drop in packed towers. Source: Eckert [15].

mathematical models representing the generalized pressure-drop curves excluding the flooding condition. Kessler and Wankat [22] formulated a model to represent Eckert's flooding curve within the range of the original data.

Operating and Design Conditions

Packed towers are typically operated at a gas velocity that corresponds to about 50-80% of flooding condition. This usually results in a ΔP of 0.5–1.0 in. H₂O/ft of packing. Alternatively, a large decrease in flow

Packir	ŋg	Cons	stants
Туре	Nominal Size, in.	α	β
Raschig rings	3/8	4.70	0.41
Ceramic	1/2	3.10	0.41
	5/8	2.35	0.26
	3/4	1.34	0.26
	1	0.97	0.25
	1 1/4	0.57	0.23
	1 1/2	0.39	0.23
	2	0.24	0.17
Raschig rings	5/8	1.20	0.28
Metal, 1/32 in.	1	0.42	0.21
Wall, 1/16 in.	1 1/2	0.29	0.20
	2	0.23	0.135
Pall rings, metal	5/8	0.43	0.17
U	1	0.15	0.16
	1 1/2	0.08	0.15
	2	0.06	0.12
Berl saddles	1/2	1.2	0.21
	3/4	0.62	0.17
	1	0.39	0.17
	1 1/2	0.21	0.13
Intalox saddles	1/2	0.82	0.20
	3/4	0.28	0.16
	1	0.31	0.16
	1 1/2	0.14	0.14

 Table 7-11

 Pressure-Drop Correlations for Packed Towers

SOURCE: Van Winkle [11].

rate causes channeling through the packed bed, and could result in poor column performance.

Two principal design methods have often been employed in sizing packed towers. One is the selection of an allowable ΔP , and the other is to select some fraction of the flooding capacity. Nguyen [23,24] uses the latter and employs Chen's least-square method to determine the tower

diameter at 50% of the superficial flooding rate. This means that the ΔPs are 2 in. H₂O/ft of bed, or greater. Here, an allowable ΔP is selected to determine the diameter of a packed tower. Kessler and Wankat's model is employed to determine the vapor rate at 70% of the flooding condition.

The maximum recommended design ΔP is 1.0 in. H₂O/ft for liquids with a specific gravity of 0.8 or more. A lower maximum design ΔP is recommended for lighter liquids. The recommended design ΔP criterion for sizing a packed tower should fall within the ranges as shown in Table 7-12 [25]. Packed beds may be scaled up using the modified Sherwood correlation as illustrated in Figure 7-14. The packing factor for various types of packing is as shown in Table 7-13 [26], because the capacity of the bed is independent of tower diameter and height. Packings that have sizes that are too large for the tower diameter will show a greater contact caused by the extra voidage, (that is, where the pieces of packing contact the tower wall) and a lower ΔP than that predicted by their published packing factors. This effect reduces to zero for most packings when the ratio of packing size to diameter exceeds 10:1 [15]. This factor is necessary when data from a pilot plant tower, which is filled with large packings, are used for scale up.

Design Equations

The packed tower sizing method is based upon the generalized ΔP correlation as shown in Figure 7-14.

Service	Design Pressure Drop in. H ₂ O/ft. of Packed Depth
Absorbers and Regenerators (Non-foaming System)	0.25-0.40
Absorbers and Regenerators	0.10-0.25
Atmospheric or Pressure Stills and Fractionators	0.40-0.80
Vacuum Stills and Fractionators	0.10-0.40

Table 7-12 Generalized Pressure Drop Rules

Table 7-13 Packing Factors

		ſ	Nominal Packing Size, in									
Type of Packing	king Mat'l.	1/4	*	1/2	%	3/4	1	1¼	1½	2	3	31/2
Super Intalox	Ceramic	-	-	-	-	-	60	-	-	30		_
Super Intalox	Plastic	-	_	-	-	-	33	-	-	21	16	
Intalox saddles	Ceramic	725	330	200	-	145	98	-	52	40	22	-
Hy-Pak rings	Metal	-	-	-	-	-	42	-	-	18	15	-
Pall rings	Plastic	-		-	97	-	52	-	40	25	-	16
Pall rings	Metai	-	-	_	70	-	48	-	28	20	-	16
Berl saddles	Ceramic	900×	-	240×	-	170 ^h	110 ^h	-	65 ^h	45×	_	-
Raschig rings	Ceramic	1,600 ^{b,×}	1,000 ^{b,×}	580°	380°	255°	155 ^d	125*.×	95°	65'	37*.×	-
Raschig rings	Metal	700×	390×	300×	170	155	115×	-	-	-		_
Raschig rings %e-in wall	Metal	-	_	410	290	220	137	110×	83	57	32×	_
Tellerettes	Plastic	-	-	_	-	-	40	-	-	20		-
Maspak	Plastic	-	-	-	-	-	-	-	-	32	20	-
Lessing exp.	Metal	-	-	-	-	-	-	-	30	-	-	-
Cross partition	Ceramic	-	-	-	-	-	-	-	-	-	70	-
• 152 Wall 552 Wall • 16 Wall 416 Wall	•%s Wall f% Wall		acking factors 16- and 30-in.		-	apolated						

Source: Eckert [26].

The abscissa on the generalized ΔP curve is expressed as:

$$X = \left(\frac{L}{G}\right) \left[\frac{\rho_{G}}{(\rho_{L} - \rho_{G})}\right]^{0.5}$$
(7-62)

where 0.01 < X < 2.0 and the ordinate is expressed as:

$$Y = \frac{(G_{I}^{2}F_{p}\mu_{L}^{0.1})}{\left[\rho_{G}(\rho_{L} - \rho_{G})g\right]}$$
(7-63)

where 0.001 < Y < 0.15

$$Y = \exp[C_0 + C_1 \ln X + C_2 (\ln X)^2 + C_3 (\ln X)^3 + C_4 (\ln X)^4]$$
(7-64)

and $C_0 - C_4$ are constants derived for each ΔP . The values of these constants for each ΔP correlation are shown in Table 7-14 [13].

The mass vapor rate, G_1 , is expressed as:

$$G_{t} = \left\{ \frac{\left[Y \rho_{G} (\rho_{L} - \rho_{G}) g \right]}{F_{p} \mu_{L}^{0.1}} \right\}^{0.5} \frac{lb}{ft^{2} \cdot s}$$
(7-65)

The tower area is

$$A = \frac{G}{(3600)(G_1)}$$
(7-66)

$$=\frac{\pi D^2}{4}$$
(7-67)

The tower diameter is defined by

$$D = \left(\frac{4A}{\pi}\right)^{0.5}, \quad \text{ft}$$
 (7-68)

At flooding condition, Eckert's flooding curve is represented as:

$$\log\left(\frac{G_2^2 F_p \Psi \mu_L^{0.2}}{\rho_G \rho_L g}\right) \quad \text{versus} \quad \log(F_{LG}) \tag{7-69}$$

		T	
Pressure	Constants	Pressure	Constants
Drop in.		Drop in.	
$H_2O/ft.$ of		${ m H_2O}/{ m ft.}$ of	
Packed Depth		Packed Depth	
0.05	$C_0 = -6.30253$	0.50	$C_0 = -4.39918$
	$C_1 = -0.60809$		$C_1 = -0.99404$
	C ₂ = -0.11932		C ₂ = -0.16983
	$C_3 = -0.00685$		C ₃ = 0.00873
	C ₄ = 0.00032		C ₄ = 0.00343
0.10	$C_0 = -5.50093$	1.0	$C_0 = -4.09505$
	$C_1 = -0.78508$		$C_{l} = -1.00120$
	$C_2 = -0.13496$		C ₂ = -0.15871
	$C_3 = 0.00134$		C ₃ = 0.00797
	$C_4 = 0.00174$		C ₄ = 0.00318
0.25	$C_0 = -5.00319$	1.5	$C_0 = -4.02555$
	$C_1 = -0.95299$		$C_1 = -0.98945$
	C ₂ = -0.13930		C ₂ = -0.08291
	C ₃ = 0.01264		C ₃ = 0.03237
	C ₄ = 0.00334		C ₄ = 0.00532

 Table 7-14

 Constants for Each Pressure Drop Correlation

Source: Blackwell [13]

where

$$F_{LG} = \left(\frac{L}{G}\right) \left(\frac{\rho_G}{\rho_L}\right)^{0.5}$$
(7-70)

Using Kessler and Wankat's model to represent the flooding condition,

$$\mathbf{X}_{1} = \mathbf{F}_{\mathrm{LG}} \tag{7-71}$$

$$Y_{1} = \frac{(G_{2}^{2}F_{p}\Psi\mu_{L}^{0.2})}{(\rho_{G},\rho_{L},g)}$$
(7-72)

At flooding condition,

$$\log(Y_1) = -1.6678 - 1.085 \log(X_1) - 0.29655 (\log X_1)^2$$
(7-73)

The mass vapor rate $lb/(sft^2)$ at 70% flooding condition can be expressed as:

$$G_{2} = 0.7 \left(\frac{Y_{1} \cdot \rho_{G} \cdot \rho_{L} \cdot g}{F_{p} \cdot \Psi \cdot \mu_{L}^{0.2}} \right)^{0.5}$$
(7-74)

The vapor velocity (ft/sec) at 70% flooding condition is expressed as:

$$V_{GF} = \frac{G_2}{\rho_G}$$
(7-75)

Guidelines for Selecting Packed Towers

Use the following rules of thumb for selecting packed towers:

- 1. Structured and random packings are suited to packed towers under 3 ft diameter and when low ΔP is desirable. In such cases, volumetric efficiences, given adequate liquid distribution throughout the column, can exceed trayed towers.
- 2. Replacing trays with packing allows greater throughput or separation in existing tower shells.
- 3. For gas rates of 500 ft³/min, use 1-inch packing. For gas rates of 200 ft³/min, or more, use 2-inch packing.
- 4. The ratio of the diameter of the tower to the packing should be at least 15.
- 5. Because of deformability, plastic packing is limited to an unsupported depth of 10–15 ft, metal to 20–25 ft.
- 6. Liquid redistributors are required every 5–10 tower diameters with pall rings, and at least every 20ft for other types of dumped packings. The number of liquid streams should be 3–5/ft² in towers larger than 3 ft diameter. Some experts advocate 9–12/ft² [27].
- 7. The height equivalent to a theoretical plate (HETP) for vaporliquid contacting is 1.3–1.8 ft for 1-inch rings, 2.5–3.0 ft for 2-inch rings.
- 8. Packed towers should operate near 70% of the flooding rate given by the correlation of Lobo *et al.* [19] and Sherwood.

9. Limit the tower height to about 175 ft maximum because of wind load and foundation considerations. Also, L/D should be less than 30.

Packed Towers Versus Trayed Towers

Both trays and packings are used to provide intimate contact between the ascending vapor and descending liquid without a great reduction in the throughput or capacity of a fractionating column. The most striking difference between trays and packings is the percentage opening of each phase-contacting device. A tray has an opening of 8–15% of the tower cross-sectional area, while the projected opening of a typical packing design is usually more than 50% of tower cross section. The void fraction of a packed column is even higher, and is usually more than 90% of the tower volume. In addition, with packing, contact is readily achieved between the vapor and the liquid phases throughout the column, rather than at specific points as with trays.

The following further illustrates the advantages of using packed towers over trays.

High capacity with high liquid rates or high viscosity. Trayed columns employ the energy of the vapor to create a mass-transfer surface area by bubbling through the liquid. Packed towers, with the aid of gravity, create a mass-transfer surface area by the action of the liquid falling over the packing. Thus, there are no downcomers in a packed tower, and 100% of the tower cross section is used for mass transfer.

High capacity/efficiency combinations. Because the capacity of a packed tower is greater than a comparable sized trayed one, a smaller, more efficient packing can be used to handle the same capacity. The range of packing sizes and types allows the combination of efficiency and capacity to be optimized.

High capacity in foaming system. Trayed columns use the continuous liquid phase to create a froth that is difficult to separate. Packed towers make the vapor phase continuous, and the liquid phase discontinuous.

Low pressure drop. Packed towers have a low ΔP per theoretical stage or transfer unit, which is beneficial in low-pressure and vacuum applications.

Low residence time. Packed towers offer low liquid holdup. This gives lower residence time for materials that are sensitive to high processing temperature. In contrast, trayed columns typically impose a holdup volume of 20–30%.

Economic Trade-Offs

In the design of industrial-scale packed towers, an economic balance is important between the capital cost of the column, its ancillary equipment, and the operating cost. If the diameter of the tower is reduced, the capacity cost will also be reduced, but the cost of pumping the gas through the column will increase because of the higher ΔP .

For columns operating at high pressures, the capital cost of the column shell is more significant, and it may be more economic to operate at gas velocities above the loading conditions. Morris and Jackson [28] suggest that a gas velocity 75–80% of the flooding velocity for normal systems, and less than 40% of the flooding rate if foaming is present.

In general, the actual operating vapor rate is some percentage of $G_{2_{nod}}$. The usual range is 65–90% of flooding, with 70-80% being most common. Because the flooding correlation is not always accurate, some confidence limit is used. For a 95% confidence, Bolles and Fair [29,30] recommend a safety factor of 1.32 for the calculated cross-sectional area. The tower diameter is calculated once the tower area is known. Because the liquid and vapor properties and gas and liquid flow rates all vary, the designer must calculate the diameter at several locations, and then use the largest value. Sizing the column diameter has often been influenced by changes in the vapor flow rate.

Eckert [21] develops an economic characterization number for various types of packing. Usually, the most suitable choice of packing is dependent on the material of construction of the equipment. In distillation towers when low-carbon steel is used, 2-inch metal rings are often favored because they give a low ΔP , high capacity, and efficiency. Alternatively, saddles are the most frequently selected plastic and ceramic packing.

Troubleshooting packed towers is well illustrated in the literature [31].

Nomenclature

A = tower area, ft^2

- $C_0 C_4$ = packed drop correlation constants
 - D = packed tower diameter, ft
 - F_{LG} = flow parameter (Equation 7-67)
 - F_p = packing factor for packing material

G = vapor rate, lb/h

 G_1 = vapor mass velocity, lb/(ft² sec tower cross section)

 G_2 = vapor mass velocity, lb/(ft² sec tower cross section)

g = gravitational constant, 32.2 ft/s^2

L = liquid rate, lb/h

 L_1 = liquid mass velocity, lb/(ft² sec tower cross section)

X = abscissa in the generalized plot

Y = ordinate in the generalized plot

 Y_1 = ordinate in Eckert's flooding curve

 V_{GF} = vapor velocity at 70% of flooding conditions, ft/s

Greek Letters

 $\alpha = \text{constant in Table 7-11}$ $\beta = \text{constant in Table 7-11}$ $\mu_L = \text{liquid viscosity, cP}$ $\rho_L = \text{liquid density, lb/ft}^3$ $\rho_G = \text{vapor density, lb/ft}^3$

 Ψ = ratio of water density to liquid density

DETERMINATION OF PLATES IN FRACTIONATING COLUMNS BY THE SMOKER EQUATIONS

Introduction

Smoker [32] developed an analytical equation that determines the number of stages when the relative volatility is constant (that is α is constant). Smoker's method can be used in preference to the McCabe-Thiele [33] graphical method for calculating the number of theoretical plates. This is because the accuracy of the McCabe-Thiele depends on the care given to the construction of the plots. The graphical construction for the number of stages becomes more difficult at very low concentrations. Also, if the relative volatility is close to one, the number of stages required will be very large, and the McCabe-Thiele diagram will be difficult to construct.

The Equations

Smoker's equations start by considering Raoult's law for a binary mixture, where α is a constant and represents the vapor-liquid equilibrium relationship. From Equation 7-25

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$$y = \frac{\alpha x}{1 + (\alpha - 1)x}$$
(7-25)

Any operating line that is either above or below the feed plate can be represented by the general equation of a straight line.

$$y = mx + b \tag{7-76}$$

Eliminating y from Equations 7-25 and 7-76 gives a quadratic equation in x.

$$m(\alpha - 1)x^{2} + [m + (\alpha - 1)b - \alpha]x + b = 0$$
(7-77)

For any particular distillation problem, Equation 7-77 will have only one real root k, between 0 and 1. Representing this root k, and substituting in Equation 7-77 gives

$$m(\alpha - 1)k^{2} + [m + (\alpha - 1)b - \alpha]k + b = 0$$
(7-78)

k is the value of the x-ordinate at the point where the extended operating lines intersect the vapor-liquid equilibrium curve.

Smoker shows that the number of stages required is given by

$$N = \log \left[\frac{x_0^* (1 - \beta x_n^*)}{x_n^* (1 - \beta x_0^*)} \right] / \log \left(\frac{\alpha}{mc^2} \right)$$
(7-79)

where

$$\beta = \frac{\mathrm{mc}(\alpha - 1)}{\alpha - \mathrm{mc}^2} \tag{7-80}$$

N = number of stages required effecting the separation represented by the concentration change from

$$x_n^*$$
 to x_0^* ; $x^* = (x - k)$ and $x_0^* > x_n^*$; $c = 1 + (\alpha - 1)k$ (7-81)

- k = composition of the liquid where the operating line intersects the equilibrium line
- m = slope of the operating line between x_n^* and x_0^*
- α = average relative volatility, assumed constant over x_n^* to $x_0^* = (\alpha_{top} + \alpha_{bottom})/2$

Application to a Distillation Column

Apply Smoker's equation to actual distillation conditions with a single feed and no side streams.

Rectifying Section:

For a rectifying column, the number of plates required to enrich from the feed composition, z_F , to the distillate (product) composition, x_D , at fixed reflux ratio, R, is required by

$$m = \frac{R}{R+1}$$
(7-82)

and

$$b = \frac{x_{\rm D}}{R+1} \tag{7-83}$$

Also

$$m(\alpha - 1)k^2 + [m + (\alpha - 1)b - \alpha]k + b = 0$$

where k is the root between 0 and 1

 $\mathbf{x}_0 = \mathbf{x}_D \tag{7-84}$

$$x_0^* = X_D - k$$
 (7-85)

$$\mathbf{x}_{n} = \mathbf{z}_{F} \tag{7-86}$$

$$x_{n}^{*} = z_{F} - k$$
 (7-87)

Stripping Section:

For the stripping section, the bottoms composition, x_B , the feed, z_F , the overhead composition, x_D and the reflux ratio, R, all fix the operating line. The values of m and b are

$$m = \frac{Rz_F + x_D - (R+1)x_B}{(R+1)(z_F - x_B)}$$
(7-88)

and

$$b = \frac{(z_F - x_D)x_B}{(R+1)(z_F - x_B)}$$
(7-89)

$$x_0^* = z_F - k$$
 (7-90)

$$x_{n}^{*} = x_{B} - k$$
 (7-91)

If the feed stream is not at its bubble point, z_F is replaced by the value of x at the intersection of operating lines given by

$$z_{\rm F}^* = \frac{b + z_{\rm F}/(q-1)}{q/(q-1) - m}$$
(7-92)

For distillation at total reflux $R = \infty$, m = 1, b = 0, k = 0, c = 1 and Equation 7-79 becomes

$$N = \log \left\{ \frac{x_0 (1 - x_n)}{x_n (1 - x_0)} \right\} / \log \alpha$$
(7-93)

Equation 7-79 simplifies to the equation derived by Underwood [34].

Smoker's equation is not limited to the range of small concentrations, but can be used for stage calculation over the entire range. In such instances, the equation is written twice, one for each section of the tower. The average relative volatilities for the rectifying section and for the stripping section are determined.

Nomenclature

- b = intercept of a straight line with the y-axis
- c = constant defined by Equation 7-81
- k = composition of the liquid where the operating line intersects the equilibrium line
- n = number of plates between any two points in the column under consideration
- R = reflux ratio, reflux per unit time divided by moles of distillate per unit time
- $x_{\rm B}$ = mole fraction in the bottoms
- $x_{\rm D}$ = mole fraction in the distillate
- z_F = mole fraction in the feed

MULTICOMPONENT DISTRIBUTION AND MINIMUM TRAYS IN DISTILLATION COLUMNS

Determining the number of stages and reflux ratio for multicomponent distillations is more complex than for binary mixtures. In multicomponent distillation with the feed containing more than two components, it is often difficult to specify the complete composition of the top and bottoms products. The separation between the top and bottoms products is specified by setting limits on two key components between which it is desired to effect separation. Also, fixing one component composition does not uniquely determine the other component compositions and the stage temperature.

Key Components

Before the start of the column design, the process designer must select the two key components between which it is desired to effect the separation. In cases where pure components are being produced, the key components are the compounds boiling adjacent to one another on the temperature scale. The material having the lower boiling point is known as the light key component and the next heavier as the heavy key component. The light key component is the component that is desired to keep out the bottoms product, and the heavy key is the component that is desired to keep out the top product. Specifications are often set on the maximum concentrations of the keys in the top and bottoms products.

The keys are known as "adjacent keys" if they are adjacent in a listing of the components in order of volatility. Sometimes, the keys selected are not adjacent but have an intermediate boiling component between them. They are known as "light key," "heavy," key and "intermediate" (boiling) or "distributed key." The non-key components that appear in both top and bottoms products are known as "distributed" components. Those that are not present to any significant extent, in one or other product, are known as the "non-distributed" components.

Equations Surveyed

The increasing availability of personal computers with acquired simulation software packages has helped design engineers to optimize a number of equilibrium stages in multipurpose fractionating towers and absorbers. Designers adept in mathematical modeling are still engaged in the use of vigorous, iterative plate-to-plate computation to study a wide range of process conditions. However, preliminary design with graphical correlations has often helped to arrive at an approximate optimum number of stages before reverting to established design methods. Figure 7-15 shows a schematic of a fractionating column with two or more multicomponent and associated equipment items. Alternatives to the preliminary design are the short-cut methods to achieve a realistic optimum number of theoretical stages. Generally, short-cut methods were proposed to establish the minimum number of stages at total reflux, and

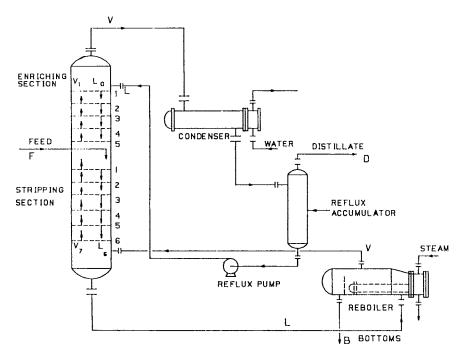


Figure 7-15. Factionation column with a reboiler and refluxing auxiliaries.

also the minimum reflux at an infinite number of theoretical stages. Many authors have correlated the minimum reflux and minimum stages with operating reflux and corresponding theoretical equilibrium stages required. These methods have enabled designers to vary the reflux ratio and plates in order to achieve an optimum relationship based on investment and operating costs.

Mathematical Modeling

The distribution of components between the distillate and bottoms is given by the Hengstebeck-Geddes equation [35,36].

$$\log\left[\frac{d_i}{b_i}\right] = A_i + B_1 \log \alpha_i$$
(7-94)

where d_i = moles of component i in the distillate

- b_i = moles of component i in the bottoms
- α_i = relative volatility of component i

 $A_1, B_1 = correlation constants$

A material balance for the ith component in the feed is:

$$\mathbf{f}_{i} = \mathbf{d}_{i} + \mathbf{b}_{i} \tag{7-95}$$

The quantity of component i in the distillate can be expressed as mole fraction recovered, or d_i/f_i . Alternatively, in the bottoms, the mole fraction of component i recovered is b_i/f_i .

If Equation 7-94 is expressed with respect to the heavy key component, then

$$\log\left[\frac{d_{HK}}{b_{HK}}\right] = A_1 + B_1 \log \alpha_{HK}$$
(7-96)

The relative volatility of the heavy key component (that is, α_{HK}) is 1.0

Therefore,

$$A_{1} = \log\left[\frac{d_{HK}}{b_{HK}}\right]$$
(7-97)

but

$$f_{HK} = d_{HK} + b_{HK} \tag{7-98}$$

and

$$\mathbf{f}_{\mathrm{LK}} = \mathbf{d}_{\mathrm{LK}} + \mathbf{b}_{\mathrm{LK}} \tag{7-99}$$

Therefore

$$A_{1} = \log\left[\frac{f_{HK} - b_{HK}}{b_{HK}}\right]$$
(7-100)

Equation 7-100 can be expressed in terms of mole fraction recovered as:

$$A_{1} = \log \left[\frac{(1.0 - b_{HK}/f_{HK})}{(b_{HK}/f_{HK})} \right]$$
(7-101)

Substituting Equation 7-97 into Equation 7-94 and expressing in terms of the light key component.

$$\log\left[\frac{d_{LK}}{b_{LK}}\right] = \log\left[\frac{d_{HK}}{b_{HK}}\right] + B_1 \log \alpha_{LK}$$
(7-102)

Therefore

$$B_{i} = \frac{\log\left[\left(\frac{d_{LK}}{b_{LK}}\right) / \left(\frac{d_{HK}}{b_{HK}}\right)\right]}{\log \alpha_{LK}}$$
(7-103)

Expressing Equation 7-103 in terms of fractional recoveries.

$$B_{1} = \frac{\log \left[\left(\frac{d_{LK}}{f_{LK}} / f_{LK}}{\frac{f_{LK}}{f_{LK}}} \right) \left(\frac{bH_{LK}}{f_{HK}} - \frac{bH_{LK}}{f_{HK}}}{f_{HK}} \right) \right]}{\log \alpha_{LK}}$$
(7-104)

$$B_{1} = \frac{\log\left[\left(\frac{d_{LK}/f_{LK}}{1 - \frac{d_{LK}}{f_{LK}}}\right)\left(\frac{bH_{LK}/f_{HK}}{1 - \frac{b_{HK}}{f_{HK}}}\right)\right]}{\log \alpha_{LK}}$$
(7-105)

The recoveries of the ith component in the distillate and bottoms:

Equation 7-94 and Equation 7-95 are

$$\log\left[\frac{d_{i}}{b_{i}}\right] = A_{1} + B_{1} \log \alpha_{i}$$

$$f_{i} = d_{i} + b_{i}$$
(7-94)
(7-95)

That is

$$\log\left[\frac{d_i/f_i}{1-d_i/f_i}\right] = A_i + B_1 \log \alpha_i$$
(7-106)

Hence,

$$\left[\frac{d_{i}/f_{i}}{1-d_{i}/f_{i}}\right] = 10^{A_{i}} \cdot 10^{(\log_{10}\alpha_{i}^{B_{i}})}$$

$$= 10^{A_{i}} \cdot \alpha_{i}^{B_{i}}$$

$$\frac{d_{i}}{1-d_{i}} = 10^{A_{i}} \cdot \alpha_{i}^{B_{i}} \left(1-\frac{d_{i}}{1-d_{i}}\right)$$
(7-108)

$$\frac{\mathbf{a}_{i}}{\mathbf{f}_{i}} = 10^{\mathbf{A}_{i}} \cdot \alpha_{i}^{\mathbf{B}_{i}} \left(1 - \frac{\mathbf{a}_{i}}{\mathbf{f}_{i}}\right)$$
(7-108)

The recovery of the ith component in the distillate is given by

$$\frac{d_{i}}{f_{i}} = \frac{(10^{A_{i}} . \alpha^{B_{i}})}{(1+10^{A_{i}} . \alpha^{B_{i}})}$$
(7-109)

And the recovery of the ith component in the bottoms is

$$\frac{b_{i}}{f_{i}} = 1 - \frac{d_{i}}{f_{i}}$$
(7-110)

$$= 1 - \frac{(10^{A_1} \cdot \alpha_i^{B_1})}{(1 + 10^{A_1} \cdot \alpha_i^{B_1})}$$
(7-111)

$$\frac{\mathbf{b}_{i}}{\mathbf{f}_{i}} = \frac{1}{1 + 10^{A_{i}} \cdot \alpha_{i}^{B_{i}}}$$
(7-112)

The Fenske's Method for Total Reflux

Fenske's [37] equation for determining the minimum equilibrium stages at total reflux was based on an ideal mixture. This suggests that the ratio of vapor pressures or the ratio of equilibrium vaporization of the key components is constant over the range of temperatures (that is, the relative volatilities are constant). Fenske expressed the minimum number of equilibrium stages as:

$$N_{\min} = \frac{\log \left[\left(\frac{X_{LK}}{X_{HK}} \right)_{D} \left(\frac{X_{HK}}{X_{LK}} \right)_{B} \right]}{\log \left[\frac{\alpha_{LK}}{\alpha_{HK}} \right]}$$
(7-113)

The Gilliland Method for Number of Equilibrium Stages

The number of theoretical equilibrium stages required for a given separation at a given reflux ratio is often determined by empirical correlations [38,39]. The abscissa, X, represents a reflux function as:

(7-114)

Correspondingly, the ordinate, Y, represents a stage function as given by

$$Y = \frac{(N - N_m)}{(N + 1)}$$
(7-115)

The ratios used for the axes of abscissa and ordinate were chosen because they provide fixed end points for the curve (X = 0.0, Y = 1.0 and at X = 1.0, Y = 0.0). The two functions were found to give good correlations. These correlations are shown in Figures 7-16 and 7-17.

Gilliland's correlation has produced relevant results that offer the following advantages:

- They represent an optimum solution concerning the location of the feed plate.
- The splitting for the two key components is verified.
- The maximum deviation using Gilliland's correlation in terms of tray number is within the 7% range [39].

The Gilliland correlation tends to be conservative for feeds with low values of the thermal condition of the feed (q). It can give inaccurate results when there is a large difference in tray requirements above and below the feed. The correlation is adequate for preliminary designs before a detailed analysis is carried out. Great caution should be exercised when used for final designs. Hines and Maddox [5] have described an improved correlation for determining the number of theoretical stages.

Underwood's Method

If an infinite or nearly infinite number of equilibrium stages is involved, a zone of constant composition must exist in the fractionating column. In this instance, there is no measurable change in the composition of liquid or vapor from stage to stage. Under these conditions, the reflux ratio can be defined as the minimum reflux ratio, R_{min} , with

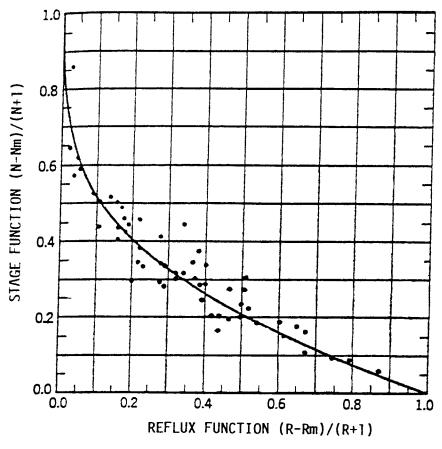


Figure 7-16. Linear coordinates based on Gilliland's correlation.

respect to a given separation of two key components (that is, the light key and heavy key) [40].

Therefore, for component i, in the distillate

$$R_{\min} + 1 = \sum_{i=1}^{n} \frac{(\alpha_i - x_{i,D})}{(\alpha_i - \theta)}$$
(7-116)

where θ is Underwood's constant (or root of the equation), and must lie between the relative volatilities of the light and heavy keys (α_{LK} and α_{HK}). The number of components is n.

Correspondingly, for the feed,

$$1 - q = \sum_{i=1}^{n} \frac{(\alpha_{i} \cdot x_{i,F})}{(\alpha_{i} - \theta)}$$
(7-117)

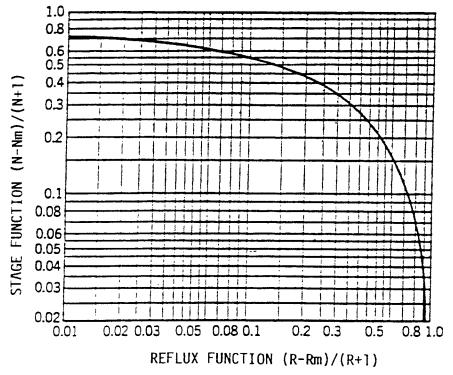


Figure 7-17. Logarithmic coordinates based on Gilliland's correlation.

Equation 7-117 shows the relationship for the feed, where q is the fraction of feed that is liquid at the feed tray temperature and pressure. For a bubble point feed, q =1, and for a dew point feed, q = 0. The minimum reflux ratio is determined from Equation 7-117 by substituting into Equation 7-116. Coker [41] developed a numerical method for computing θ and R_{min} respectively. However, other methods should be tried, if R_{min} gives a negative value. Also, it may be that the separation between the feed and the overhead can be accomplished in less than one equilibrium stage.

Equations for Describing Gilliland's Graph

Many equations have been proposed to describe Gilliland's curve for multicomponent distillation. However, the difficulty with some of these equations has been in meeting the end conditions of X = 0, Y = 1 and X = 1, Y = 0. A review of the many equations proposed by these authors is as follows:

Proposed Equations:

1. Hengstebeck [42] (1961) $\log Y = A + B(\log X) + C(\log X)^{2} + D(\log X)^{3} + E(\log X)^{4}$ (7-118) where A = -1.3640187B = -3.0920489C = -3.407344729D = -1.74673876E = -0.332688972. Liddle [43] (1968) For $0.0 \le X \le 0.01$, Y = 1.0 - 18.5715X(7-119)For $0.01 \le X \le 0.90$, Y = 0.545827 - 0.591422X+ 0.002743/X(7-120)For $0.90 \le X \le 1.0$, Y = 0.16595 - 0.16595X(7-121)3. Van-Winkle and Todd [44] (1971) For 0.0078 < X < 0.125, Y = 0.5039 - 0.5968X $-0.0908(\log X)$ (7-122)For 1.25 < X to X = 1.0, $Y = 0.6257 - 0.9868X + 0.516X^2$ $-0.1738X^{3}$ (7-123)4. Molokavov [45] (1972) $0 \le (X, Y) \le 1.0$ $Y = 1 - \exp\left[\left(\frac{1 + 54.4X}{11 + 117.2X}\right)\left(\frac{X - 1}{X^{0.5}}\right)\right]$ (7-124)5. Hohman and Lockhart [46] (1972) X = 0, Y = 0.65X = 1.0, Y = 0.067 $Y = \frac{0.65 - 0.50X}{1.0 + 1.25X}$ (7-125)

6. Eduljee [47] (1975)

$$X = 0, Y = 1.0$$

 $X = 1.0, Y = 0$
 $Y = 0.75 - 0.75X^{0.5668}$ (7-126)
7. Huan-Yang Chang [48] (1981)
For $X = 0, Y = 1.0$
 $X = 1.0, Y = 0$
 $Y = 1 - \exp\left(1.49 + 0.315X - \frac{1.805}{X^{0.1}}\right)$ (7-127)
8. Harg [49] (1985)
For $X = 0, Y = 1.0$
 $X = 1.0, Y = 0$
 $Y = 1 - X^{1/3}$ (7-128)
9. McCormick [50] (1988)
 $X = 0, Y = 1.0$
 $X = 1.0, Y = 0$
 $Y = 1 - X^B$

where

$$B = 0.105(\log X) + 0.44 \tag{7-129}$$

From the equations listed, McCormick's gives a good agreement in the normal operating range of real towers. The reflux ratio, R, is calculated as a multiple of the minimum reflux ratio, R_{min} .

That is,

$$R = FACTOR.R_{min}$$
(7-130)

The multiplier "FACTOR" generally varies from 1.2 to about 1.5 for conventional columns, but because of economic situations, the range is now between 1.05 and 1.20.

Kirkbride's Feed Plate Location

After the minimum number of stages and the minimum reflux ratio have been determined, the number of theoretical stages is then calculated. The ratio of the number of plates above the feed stage (including the partial condenser) to the number below the feed stage (including the reboiler) can be obtained using Kirkbride's empirical equation [51]. The equation was developed on the basis that the ratio of rectifying trays to stripping trays depends on:

- The fraction of the heavy key component (in the feed) removed in the overhead.
- The fraction of the light key component removed in the bottoms.
- The concentration of the heavy key component in the overhead.
- The concentration of the light key component in the bottoms.

Kirkbride's feed plate equation is expressed as:

$$\log\left[\frac{m}{p}\right] = 0.206 \log\left\{\left(\frac{B}{D}\right)\left(\frac{x_{HK}}{x_{LK}}\right)_{F}\left[\frac{(x_{LK})_{B}}{(x_{HK})_{D}}\right]^{2}\right\}$$
(7-131)

where B = molar flow bottoms product

- D = molar flow top product
- m = number of theoretical stages above the feed plate, including any partial condenser
- p = number of theoretical stages below the feed plate, including the reboiler
- $(x_{LK})_F$ = concentration of the light key in the feed
- $(x_{HK})_{F}$ = concentration of the heavy key in the feed
- $(x_{LK})_B$ = concentration of the light key in the bottoms product
- $(x_{HK})_{D}$ = concentration of the heavy key in the top product

Nomenclature

 A_1, B_1 = correlation constants in Hengstebeck-Geddes equation

- b = number of moles of a component in the bottoms
- B = bottoms flow rate moles/hr
- d = number of moles of a component in the distillate
- D = distillate flow rate moles/hr
- F = feed flow rate moles/hr
- f = number of moles of a component in the feed

FACTOR = multiplier for the reflux ratio (that is, FACTOR R_{min})

K = equilibrium constant for a component in the feed

n = number of components

- N = number of theoretical plates
- N_{min} = minimum number of theoretical plates at total reflux
 - q = pseudo-ratio of liquid to vapor for the feed

R = reflux ratio

 R_{min} = minimum reflux ratio

x = mole fraction of a component where $x_B, x_D, x_F =$ mole fractions in the bottoms, distillate, and feed, respectively

Greek Letters

- α = relative volatility with respect to the heavy key
- θ = Underwood's constant
- $\Sigma =$ summation

HK, LK = heavy and the light key components

i = individual component identifier

PROBLEMS AND SOLUTIONS

Problem 7-1

A multicomponent hydrocarbon gas mixture is composed as follows:

Component	Mol %	к
Methane (CH_4)	27.52	7.88
Ethane (C_2H_6)	16.34	2.77
Propane (C_3H_8)	29.18	1.18
Isobutane (iC_4H_{10})	5.37	0.61
Normal butane (nC_4H_{10})	17.18	0.48
Isopentane (iC_5H_{12})	1.72	0.264
Normal pentane(nC_5H_{12})	2.18	0.225
Hexane (C_6H_{14})	0.47	0.102
Heptane (C ₇ H ₁₆)	0.04	0.048

Operating temperature and pressure are 178°C and 400 psia. Determine the dew point temperature of the gas mixture.

Solution

The computer program PROG71 calculates the dew and bubble points of any multicomponent hydrocarbon mixture based on the user supplied K-values. The program will handle feed streams containing up to 15 components. The feed entries may be moles, mole fraction, or mole percent. The dew point ($\Sigma X = \Sigma Y/K$) should be 1.0 or very close to 1.0, depending on the tolerance the user will accept. If the sum printed is not equal to 1.0 (within the user tolerance), a message is printed instructing the user to assume a new temperature. The iteration is repeated using the K_i's corresponding to the new temperature chosen. The temperature supplied does not affect the computation, but is printed and stored as a reminder to the user of which temperature (and corresponding K-values) was used for the last iteration. Table 7-15 shows the input data and computer output instructing the user that the dew point temperature is increased.

At 200 psia,	, calculate th	he bubble	e point	temperature	of the foll	owing:

Component	Mol %	K @ 260°F	K @ 235°F	K @ 237°F
iC ₄ H ₁₀	18.2	1.92	1.62	1.65
nC ₄ H ₁₀	23.8	1.58	1.35	1.35
iC ₅ H ₁₂	33.7	0.93	0.76	0.77
nC_5H_{12}	12.1	0.81	0.64	0.64
C ₆ H ₁₄	12.2	0.42	0.315	0.32

Solution

The computer program PROG71 determines the bubble points of the hydrocarbon mixture. As with the dew point, the bubble point $(\Sigma Y = \Sigma KX)$ is computed until the sum is 1.0 or very close to 1.0, depending on the tolerance the user will accept. If the sum printed is not equal to 1.0 (within tolerance), the iteration is repeated using the K_i's corresponding to the new temperature. Tables 7-16, 7-17, and 7-18 give both the data input and computer outputs for the bubble points of the hydrocarbon mixture at 260°F, 235°F and 237°F.

Problem 7-2

The composition of the feed to a natural gas liquefaction plant is shown in the feed stream data. The feed will be flashed at 600 psia and

Table 7-15
Input Data and Computer Output for Dew
and Bubble Points Calculations

EW			
78			
7.52	7.88		
	2.77		
9.18	1.18		
.37	0.61		
7.18	0.48		
.72	0.264		
.18			
	0.102		
.04	0.048		
	DEW AND	BUBBLE POINTS CALCULATIONS	
*****	*****	********	*****
D	EW POINT CALCULA	TION	
т	EMPERATURE OF: 17	8.00	
N	UMBER OF COMPONENTS	IN THE FEED STREAM: 9	
N	UMBER OF COMPONENTS	FEED COMPOSITION (MOLE %)	K-VALUE
	1	27.5200	7.8800
	2	16.3400	2.7700
	3 4	29.1800	1.1800
		5.3700 17.1800	.6100 .4800
	E		
	5		
	6	1.7200	.2640
	6 7	1.7200 2.1800	•2640 •2250
	6	1.7200	.2640
 T	6 7 8	1.7200 2.1800 .4700	.2640 .2250 .1020
	6 7 8 9	1.7200 2.1800 .4700 .0400	.2640 .2250 .1020
	6 7 8 9 OTAL COMPOSITION: UMBER OF COMPONENTS 1	1.7200 2.1800 .4700 .0400 100.0000 MOLE FRACTION .2752	.2640 .2250 .1020 .0480 X=Y/K .0349
	6 7 8 9 OTAL COMPOSITION: UMBER OF COMPONENTS 1 2	1.7200 2.1800 .4700 .0400 100.0000 MOLE FRACTION .2752 .1634	.2640 .2250 .1020 .0480 X=Y/K .0349 .0590
	6 7 8 9 OTAL COMPOSITION: UMBER OF COMPONENTS 1 2 3	1.7200 2.1800 .4700 .0400 100.0000 MOLE FRACTION .2752 .1634 .2918	.2640 .2250 .1020 .0480 .0480 .0349 .0590 .2473
	6 7 8 9 OTAL COMPOSITION: UMBER OF COMPONENTS 1 2 3 4	1.7200 2.1800 .4700 .0400 100.0000 MOLE FRACTION .2752 .1634 .2918 .0537	.2640 .2250 .1020 .0480 X=Y/K .0349 .0590 .2473 .0880
	6 7 8 9 OTAL COMPOSITION: UMBER OF COMPONENTS 1 2 3 4 5	1.7200 2.1800 .4700 .0400 100.0000 MOLE FRACTION .2752 .1634 .2918 .0537 .1718	.2640 .2250 .1020 .0480 .0480 .0349 .0590 .2473 .0880 .3579
	6 7 8 9 OTAL COMPOSITION: UMBER OF COMPONENTS 1 2 3 4 5 6	1.7200 2.1800 .4700 .0400 100.0000 MOLE FRACTION .2752 .1634 .2918 .0537 .1718 .0172	.2640 .2250 .1020 .0480 .0480 .0480 .0590 .2473 .0880 .3579 .0652
	6 7 8 9 OTAL COMPOSITION: UMBER OF COMPONENTS 1 2 3 4 5 6 7	1.7200 2.1800 .4700 .0400 100.0000 MOLE FRACTION .2752 .1634 .2918 .0537 .1718 .0172 .0218	.2640 .2250 .1020 .0480 .0480 .0590 .2473 .0880 .3579 .0652 .0969
	6 7 8 9 OTAL COMPOSITION: UMBER OF COMPONENTS 1 2 3 4 5 6 7 8	1.7200 2.1800 .4700 .0400 100.0000 MOLE FRACTION .2752 .1634 .2918 .0537 .1718 .0172 .0218 .0047	.2640 .2250 .1020 .0480 .0480 .0590 .2473 .0880 .3579 .0652 .0969 .0461
N 	6 7 8 9 OTAL COMPOSITION: UMBER OF COMPONENTS 1 2 3 4 5 6 7	1.7200 2.1800 .4700 .0400 100.0000 MOLE FRACTION .2752 .1634 .2918 .0537 .1718 .0172 .0218	.2640 .2250 .1020 .0480 .0480 .0349 .0590 .2473 .0880 .3579 .0652 .0969

RAISE THE DEW-POINT TEMPERATURE: OF

	······································	
DATA71.DAT		
BUBBLE		
260		
5		
8.2 1.92		
23.8 1.58 33.7 0.93		
2.1 0.81		
2.2 0.42		
	BUBBLE POINTS CALCULATIONS	
	BUBBLE POINTS CALCULATIONS	*****
BUBBLE POINT CALCULAT	ION	
TEMPERATURE oF: 260	.00	
NUMBER OF COMPONENTS I	N THE FEED STREAM: 5	
NUMBER OF COMPONENTS	FEED COMPOSITION (MOLE %)	K-VALUE
1	18.2000	1.9200
2	23.8000	1.5800
3	33.7000	.9300
4	12.1000	.8100
5	12.2000	.4200
TOTAL COMPOSITION:	100.0000	
NUMBER OF COMPONENTS	MOLE FRACTION	Y=KX
1	.1820	.3494
2	.2380	.3760
3	.3370	.3134
4	.1210	.0980
5	.1220	.0512
		1
TOTAL BUBBLE POINT:		1.1881

Table 7-16 Input Data and Computer Output for Dew and Bubble Points Calculations

LOWER THE BUBBLE POINT TEMPERATURE: OF

20°F. If the flow rate is 1,000 moles/h, calculate the flow rates of the liquid and vapor streams and their compositions.

Feed stream data:

Number	Component Name	Feed Mole Fraction	Equilibrium Constant, K
1	CO ₂	0.0112	0.90
2	CH_4	0.8957	2.70
3	C_2H_6	0.0526	0.38
4	C_3H_8	0.0197	0.098
5	iC ₄ H ₁₀	0.0068	0.038

(table continued on next page)

Number	Component Name	Feed Mole Fraction	Equilibrium Constant, K
6	nC_4H_{10}	0.0047	0.024
7	$C_{5}H_{12}$	0.0038	0.0075
8	$C_{6}H_{14}$	0.0031	0.00019
9	$C_7 H_{16}$ and heavier	0.0024	0.0007

Table 7-16 (continued)

Solution

The computer program PROG72 performs equilibrium flash calculations for an ideal multi-component mixture. The program listed determines the moles of each component in the liquid and vapor phases

DATA71.DAT -----BUBBLE 235
 18.2
 1.62

 23.8
 1.35

 33.7
 0.76

 12.1
 0.64

 12.2
 0.315
 DEW AND BUBBLE POINTS CALCULATIONS ***** BUBBLE POINT CALCULATION TEMPERATURE oF: 235.00 NUMBER OF COMPONENTS IN THE FEED STREAM: 5 FEED COMPOSITION (MOLE %) K-VALUE NUMBER OF COMPONENTS _____ ~~-----18.2000 1.6200 1 1.3500 2 23.8000 .7600 3 33,7000 .6400 4 12.1000 5 12.2000 .3150 _____ TOTAL COMPOSITION: 100.0000 NUMBER OF COMPONENTS MOLE FRACTION Y=KX _____ ------------.1820 .2948 1 .2380 .3213 2 .3370 3 .2561 .0774 4 .0384 .1220 5 _____ _____ TOTAL BUBBLE POINT: .9881

Table 7-17 Input Data and Computer Output for Dew and Bubble Points Calculations

RAISE THE BUBBLE POINT TEMPERATURE: oF

DATA71.DAT						
BUBBLE						
235						
5						
1.65						
23.8 1.35 33.7 0.77						
12.1 0.64						
12.2 0.32						
DEW AND BUBBLE POINTS CALCULATIONS						
BUBBLE POINT CALCULAT	EON					
TEMPERATURE oF: 235	.00					
NUMBER OF COMPONENTS IN	N THE FEED STREAM: 5					
NUMBER OF COMPONENTS	FEED COMPOSITION (MOLE %)	K-VALUE				
	18.2000	1.6500				
1	18.2000	1.0300				
1 2	23.8000	1.3500				
2 3	23.8000 33.7000	1.3500 .7700				
2 3 4	23.8000 33.7000 12.1000	1.3500 .7700 .6400				
2 3	23.8000 33.7000 12.1000 12.2000	1.3500 .7700				
2 3 4	23.8000 33.7000 12.1000	1.3500 .7700 .6400				
2 3 4 5 TOTAL COMPOSITION:	23.8000 33.7000 12.1000 12.2000	1.3500 .7700 .6400				
2 3 4 5 TOTAL COMPOSITION:	23.8000 33.7000 12.1000 12.2000 100.0000	1.3500 .7700 .6400 .3200 Y=KX .3003				
2 3 4 5 TOTAL COMPOSITION: NUMBER OF COMPONENTS 1 2	23.8000 33.7000 12.1000 12.2000 100.0000 MOLE FRACTION .1820 .2380	1.3500 .7700 .6400 .3200 				
2 3 4 5 TOTAL COMPOSITION: NUMBER OF COMPONENTS 1 2 3	23.8000 33.7000 12.1000 12.2000 100.0000 MOLE FRACTION .1820 .2380 .3370	1.3500 .7700 .6400 .3200 Y=KX .3003 .3213 .2595				
2 3 4 5 TOTAL COMPOSITION: NUMBER OF COMPONENTS 1 2 3 4	23.8000 33.7000 12.1000 12.2000 100.0000 MOLE FRACTION .1820 .2380 .3370 .1210	1.3500 .7700 .6400 .3200 Y=KX .3003 .3213 .2595 .0774				
2 3 4 5 TOTAL COMPOSITION: NUMBER OF COMPONENTS 1 2 3	23.8000 33.7000 12.1000 12.2000 100.0000 MOLE FRACTION .1820 .2380 .3370	1.3500 .7700 .6400 .3200 Y=KX .3003 .3213 .2595				

Table 7-18 Input Data and Computer Output for Dew and Bubble Points Calculations

including the feed fraction condensed or vaporized. In addition, all feed moles and K-values are printed. As an added feature, the program checks the feed composition at the conditions given. This determines whether the feed is all vapor or all liquid. These computations are performed before the flash calculations are started, and a message is printed if either condition exists. Table 7-19 gives the input data and the computer output of the multicomponent mixture.

Problem 7-3*

Size an absorber tower for a methy ethyl amine (MEA) system with the following conditions:

Table 7-19 Input Data and Computer Output for Multicomponent Equilibrium Flash Calculations

DATA72.DAT					
20.0					
600.0					
9					
11.2	0.90				
895.7	2.70				
52.6	0.38				
19.7	0.098				
6.8	0.038				
4.7	0.024				
3.8	0.0075				
3.1	0.00019				
2.4	0.0007				

MULTICOMPONENT EQUILIBRIUM FLASH CALCULATIONS

MULTICOMPONENT EQUILIBRIUM FLASH CALCULATION AT 20.0 DEG. F AND 600.0 psia

COMPO	DNENT K-VA	ALUE FE	ED	LIQU	ID	VAI	OR
NUMBE	ER	Mols/h.	Mol. frac.	Mols/h.	Mol. frac.	Mols/h.	Mol. frac.
1	.900	11.200	.011	.510	.012	10.690	.011
2	2.700	895.700	.896	14.030	.341	881.670	.920
3	.380	52.600	.053	5.343	.130	47,257	.049
4	.098	19.700	.020	6.004	.146	13.696	.014
5	.038	6.800	.007	3.608	.088	3.192	.003
6	.024	4.700	.005	3.016	.073	1.684	.002
7	.007	3.800	.004	3.235	.079	.565	.001
8	.000	3.100	.003	3.086	.075	.014	.000
9	.001	2.400	.002	2.362	.057	.038	.000
TOT	ALS	1000.000	1.000	41.195	1.000	958.805	1.000

Vapor flow	40,000 lb/h.
Vapor density	0.295 lb/ft3
Liquid flow	330,000 lb/h.
Liquid density	61.85 lb/ft ³
Foam factor	0.75
Residence Time	4.5s
Tray Spacing	24in.
Surface tension	57.6 dyne/cm.

The author developed computer program PROG73 for solving problem 7-3 in the text, using Nutter's *Float Valve Design Manual*. The developed program and calculation of the tray geometry are the responsibility of the author. The author expresses his gratitude to Nutter Engineering for the use of the photographs and the information from the Float Valve Design Manual.

Solution

The computer program PROG73 determines the parameters required for sizing a tower using valve trays. The program calculates the bubbling area, downcomer area, tower area, and diameter. The tower diameter may be rounded to an appropriate value and the area is recalculated. Once the tower diameter is set, the downcomer area ratio is determined and the downcomer height is computed from Tables 7-5 to 7-9. Table 7-20 shows the input data and computer output for tower sizing using valve trays. The computer output gives the tower area of 26.571 ft² and a diameter of 5.816 ft.

Tray Geometry

From the tower size (normally by increasing the diameter to the nearest 6-inch increment), the following procedure is used to calculate the downcomer dimensions.

DATA73.D	AT	
	330000.0 61.85 0.75 4.5 57.6	
	TOWER DESIGN FOR VALVE TH	
VAPO VAPO LIQU LIQU VAPO FOAM DOWN TRAY SURF SAFE DENS BUBB DOWN TOWE TOWE	R FLOW RATE, lb/hr.: R VOLUMETRIC RATE, ft^3/sec.: ID FLOW RATE, lb/hr.: ID VOLUMETRIC RATE, ft^3/sec.: ID DENSITY, lb/ft^3.: FACTOR: COMER RESIDENCE TIME, sec.: SPACING, in.: ACE TENSION, dyne/cm.: ACE TENSION, dyne/cm.: ACE TENSION/VAPOR DENSITY (X): TY FACTOR: ITY RADICAL: LING AREA, ft^2: COMER AREA, ft^2: R AREA, ft^2: R DIAMETER, ft: COMER AREA RATIO:	40000.0 37.665 330000.0 1.482 61.850 .295 .750 4.500 24. 57.60 195.254 .745 .069 10.905 4.446 26.571 5.816 .2246

Table 7-20 Input Data and Computer Output for Tower Design Using Valve Trays

The downcomer area ratio is:

$$\mathbf{R}_{\mathrm{DCA}} = \frac{\mathbf{A}_{\mathrm{DC}}}{(2\mathbf{A}_{\mathrm{DC}} + \mathbf{A}_{\mathrm{b}})}$$

The calculated tower diameter is increased to 6.0 ft and the tower area becomes

$$A = \frac{\pi \cdot d^2}{4}$$
$$= \frac{\pi \times 6^2}{4}$$
$$= 28.27 \text{ ft}^2$$

Using Tables 7-5 to 7-9 and R_{DCA} will allow a direct computation of chord height for the side downcomer. This is rounded up to the nearest 1/2-inch incremental dimension. Using this dimension, the corrected downcomer area is calculated from Tables 7-5 to 7-9. The computed output of the tower used by the downcomer area (R_{DCA}) is 0.2246. That is,

$$\frac{A_d}{A_t} = 0.2246$$

	L DIA	$\frac{A_{d}}{A_{t}}$
0.2760	0.8940	0.2246

The corresponding downcomer height-to-tower-diameter ratio is 0.2760. The downcomer height is (0.2760) (6) (12)

$$= 19.87 \text{ in.} + 1/2 \text{ in.}$$

= 20.4 in.

The new
$$\frac{H}{DIA} = \frac{20.4}{72}$$

= 0.2833

This gives a downcomer-area-to-tower-area ratio of 0.2329.

	L DIA	$\frac{A_{d}}{A_{t}}$
0.2830	0.9009	0.2326
Avg. 0.28325	Avg. 0.90115	Avg. 0.2329
0.2835	0.9014	0.2332

That is

$$\frac{A_d}{A_t} = 0.2329$$

The downcomer area, A_d , is (0.2329) (28.27) = 6.584 ft²

The weir length, L, is (0.90115×72) = 64.88 in. = 65 in.

The computed liquid flow rate is 1.482 ft³/sec or 664.907 USgpm.

conversion: 1 ft³/sec = 448.655 USgpm

The liquid flow per inch of weir = $\frac{664.907}{65}$

= 10.23 USgpm per inch of weir

A single-pass tray is selected.

Problem 7-4

A packed tower with 1-inch Raschig rings are to be used to absorb ammonia from air by contacting the gas stream with water. The entering gas flow is 40 lb mol/h and contains 5.0% NH₃. Ninety percent of the NH₃ is to be removed from the air. The entering water flow rate is 3,200 lb/h. Absorption is to be carried out at 1 atm. abs. and 20°C. Estimate:

1. The diameter of the tower for a ΔP of 1.0 in. H₂O/ft. packing.

2. The vapor rate at 70% of flooding condition.

Physical properties:

$$\rho_{\rm G} = (28.4) \left(\frac{1}{359}\right) \left(\frac{273}{293}\right)$$

= 0.0737 lb/ft³
$$\rho_{\rm L} = 62.3 \, \text{lb/ft}^3$$

$$\mu_{\rm L} = 1 \text{cP}$$

$$F_{\rm p} = 155$$

Solution

When calculating the tower diameter, conditions at the bottom of the tower should be used because this corresponds to the maximum flow rates. At the bottom of the column:

$$MW_{gas} = 0.05 \times 17 + 0.95 \times 29$$

= 28.4 lb/lb. mol.

Molecular weight of ammonia $\left(MW_{NH_3} = 17 \frac{lb}{lb. mol}\right)$

 NH_3 removed from the air = (0.9)(0.05)(40)

$$= \left(1.8 \frac{\text{lb.mol}}{\text{hr}}\right) \left(17 \frac{\text{lb}}{\text{lb.mol}}\right)$$
$$= 30.6 \text{ lb/hr}$$

Gas flow rate at the bottom = $\left(40 \frac{\text{lb.mol}}{\text{hr}}\right) \left(28.4 \frac{\text{lb}}{\text{lb.mol}}\right)$

= 1,136 lb/hr

Liquid flow rate at the bottom = 3200 + 30.6

The computer program PROG74 calculates the diameter of a packed tower for pressure drops of 0.05, 0.10, 0.25, 0.50, 1.0 and 1.5 in H_2O/ft .

The simulation is based on the generalized pressure drop correlation of Sherwood *et al.* [20].

The curves have been fitted to a series of equations developed by Blackwell [13]. In addition, the program computes the vapor rate and vapor velocity at 70% of flooding condition. Table 7-21 illustrates the input data and computer output of the packed tower.

Problem 7-5

A column is to be designed to separate 1000 moles/hr of a binary mixture of benzene (C_6H_6) and toluene (C_7H_8). The feed will contain 40% benzene and 60% toluene. A distillate that is 99% benzene and a bottoms that is 1% benzene are desired at a reflux ratio of 3 to 1. For this mixture, the average value of relative volatility (α) is 2.50. Estimate the number of equilibrium stages at this reflux ratio and the optimum feed stage location.

Solution

The computer program PROG75 uses the analytical method of a quadratic equation to determine the number of stages in the rectifying and stripping sections of a distillation column. The program uses Smoker's equations involving two components to calculate the required number of stages. Table 7-22 gives the input data and computer output. The

 Table 7-21

 Input Data and Computer Output for Packed Tower Design

DATA74.DAT					
3230.6	1136.0 155.0	0.073	7		
			PACKED TOWER DESIGN	*****	
******	UID FLOW	። የ እስጥድ	lb/hr.:	3230.600	
	FLOW RA			1136.000	
	DENSITY			.0737	
				62.3000	
LIQUID DENSITY, lb/ft.^3: TOWER PACKING FACTOR:				155.0	
PRESSURE DROP, INCH-H2O:				1.000	
LIQUID VISCOSITY, CP:				1.000	
			THE GENERALIZED PLOT:	.0979	
			THE GENERALIZED PLOT:	.0719	
			RATE, lb/s.ft^2:	.2617	
			FLOODING CONDITION:	.0978	
YC	O-ORDINA	TE AT	FLOODING CONDITION:	.1335	
VAF	OR RATE	FLOODING CONDITION, 1b/s.ft^2:	.2496		
VAPOR VELOCITY AT 70% FLOODING CONDITION, ft./sec:				3.39	
	ER AREA			1.21	
TOP	ER DIAM	ETER, i		14.87	
TOP	ER AREA	AT 70%	FLOODING CONDITION, ft^2:	1.67	
TOP	ER DIAM	ETER AT	70% FLOODING CONDITION, inches:	17.49	

 Table 7-22

 Input Data and Computer Output for Calculating Equilibrium

 Number of Stages Using Smoker's Equations

DATA75.DAT		
2.5 2.5 0.40 0.99 0.01 3		
EQUILIBRIUM NUMBER OF STAGES BY SMOKER'S ************************************	EQUATION 2.500 2.500	
THE MOLE FRACTION OF THE FEED: THE MOLE FRACTION OF THE DISTILLATE: THE MOLE FRACTION OF THE BOTTOMS: THE REFLUX RATIO:	.4000 .9900 .0100 3.00	
THE NUMBER OF STAGES IN THE RECTIFYING SECTION: THE NUMBER OF STAGES IN THE STRIPPING SECTION: TOTAL NUMBER OF STAGES: FEED STAGE LOCATION:	7.87 7.76 15.63 7.8	

required number of stages between benzene and toluene mixture is 16. The optimum feed location is 8.

Problem 7-6

The feed to a butane-pentane splitter of the following composition is to be fractionated into a distillate product that contains 95% of the n-butane contained in the feed. The bottoms product contains 95% of the iso-pentane in the feed. The reflux ratio for the fractionation will be 1.3 R_{min} , and the column pressure will be 100 psia at the top plate. The reflux and feed are at their bubble point temperatures. The feed composition and equilibrium constant are:

Number	Component	X _F	K _i
1	iC_4H_{10}	0.06 ,	2.15
2	$nC_4H_{10}(LK)$	0.17	1.70
3	iC ₅ H ₁₂ (HK)	0.32	0.835
4	nC ₅ H ₁₂	0.45	0.70

Use the short-cut method to determine:

- 1. The recoveries of the components in the distillate and bottoms.
- 2. The minimum number of stages.
- 3. The minimum reflux ratio.
- 4. The actual number of theoretical plates.
- 5. The optimum location of the feed plate.

Solution

The computer program PROG76 gives the multicomponent distribution calculations based on a series of equations developed by Hengstebeck-Geddes [35,36], while the minimum tray calculations (at total reflux) use the equations developed by Fenske [37]. The number of theoretical equilibrium stages is determined using Gilliland's [38,39] empirical correlations. The program listed calculates the following: feed component recovery in both the distillate and bottoms products, the minimum number of stages, Underwood constant, minimum reflux ratio, actual reflux ratio, total number of theoretical plates, and the optimum location of the feed plate in a distillation column. The user has an option of entering the feed components (moles) and the equilibrium constants or the feed and the relative volatilities. Feed components must be entered in order of decreasing relative volatility, with the light key (LK) and heavy key (HK) components adjacent to each other. Table 7-23 shows the input data and computer output of the butane-pentane splitter.

 Table 7-23

 Input Data and Computer Output for Multicomponent

 System Fractionation

DATA76.DA	T						
EQUIL							
4							
0.06 2							
0.17 1							
0.45 0	.70						
0.45 C	.95						
3 1 1	3						
		MULTICOMPON	NENT SY	STEM FI	RACTIONATIO	N	
*******	*****	*********	******	*****	*********	********	*******
COMPONENT	FEED	REL VOLATI	ILITY	DISTI	LLATE	BOTTOMS	
NUMBER	MOLES	ALPHA		*	MOLES	*	MOLES
1	.0600 .1700 .3200	2.5749	99.2	532	.0596	.7468	,0004
2	.1700	2.0359	95.0	000	.1615	5.0000	.0085
3	.3200	1.0000	5.0	000	.0160	95.0000	.3040
4	.4500	.8383	1.2	067	.0054	98.7933	.4446
THE HEA	VY KEY COMPONE	NT NUMBER:				3	
	AGE RECOVERY O		KEY CC	MPONEN	г		
	DISTILLATE (%)					95.00	
	AGE RECOVERY O	F THE HEAVY	KEY CO	MPONEN	r	95.00	
	BOTTOMS (%):	· .				8.3	
MINIMUM NUMBER OF STAGES: TOTAL MOLES IN THE DISTILLATE:				.2425			
TOTAL MOLES IN THE BOTTOMS:				.7575			
UNDERWOOD CONSTANT:				1.6213			
MINIMUM REFLUX RATIO:				2.8036			
ACTUAL REFLUX RATIO:				3.6447			
NUMBER OF THEORETICAL PLATES IN THE COLUMN:			16.2				
THE POS	SITION OF THE F	EED PLATE:				6.7	

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PROGRAM PROG71

```
CALCULATION OF THE DEW POINT AND BUBBLE POINT OF GAS MIXTURE.
С
С
         THIS PROGRAM WILL CALCULATE DEW POINT AND BUBBLE POINT BASED
с
         ON THE SUPPLIED K-VALUES.
00000000
         THE PROGRAM WILL COVER FEED STREAMS CONTAINING UP TO 15
         COMPONENTS.
         Ki = EQUILIBRIUM FLASH CONSTANT FOR EACH FEED STREAM
         MI = TOTAL MOLES OF COMPONENT I IN THE FEED STREAM
         NI = MOLE FRACTION OF COMPONENT I IN THE FEED STREAM
                                                                      **********
         CHARACTER*10 FEED
с
         COMMON/DATA/BP, DP, J, K, N
         REAL K(20), M(20), N(20), BP(20), DP(20)
         OPEN (UNIT = 3, FILE='DATA71.DAT', STATUS='OLD', ERR=18)
         OPEN (UNIT = 1, FILE='PRN')
С
С
         READ EITHER THE DEW POINT OR BUBBLE POINT TEMPERATURE
С
         READ THE TOTAL NUMBER OF COMPONENTS
         READ THE MOLES OF EACH FEED COMPONENT
READ THE K-VALUE OF EACH COMPONENT
С
с
с
         READ (3, 10, ERR=19) FEED
         FORMAT (A6)
 10
         READ (3,*,ERR=19) T
READ (3,*,ERR=19) J
         READ (3,*,ERR=19)(M(I),K(I),I=1,J)
         WRITE (1, 100)
        FORMAT (///,20x,'DEW AND BUBBLE POINTS CALCULATIONS',/1H ,
76(1H*))
 100
         WRITE (1, 110) FEED
         FORMAT (/,6X, A6, 2X, 'POINT CALCULATION')
WRITE (1, 120) T
FORMAT (/,6X,'TEMPERATURE oF:',F10.2)
 110
 120
         WRITE (1, 130) J
FORMAT (/,6X,'NUMBER OF COMPONENTS IN THE FEED STREAM:',12)
 130
 130 FORMAT (/,0.4, BORDER OF COMPONENTS',6X,'FEED COMPOSITION (MOLE %)',
WRITE (1, 140)
140 FORMAT (/,6X,'NUMBER OF COMPONENTS',6X,'FEED COMPOSITION (MOLE %)',
* 6X,'K-VALUE',/1H ,76(1H-))
         DO I = 1, J
WRITE (1, 150) I,M(I),K(I)
         FORMAT (13X, 12, 24X, F8.4, 16X, F8.4)
 150
         END DO
         GO TO 20
         WRITE (1, 160)
   18
         FORMAT (6X, 'DATA FILE DOES NOT EXIST')
 160
         GO TO 999
```

```
19
      WRITE (1, 170)
      FORMAT (6X, 'ERROR MESSAGE IN THE DATA VALUE')
170
      GO TO 999
20
      SUM=0.0
      DO I = 1, J
      SUM=SUM+M(I)
      END DO
      WRITE (1, 180)
      FORMAT (1H ,76(1H-))
180
      WRITE (1, 190)SUM
      FORMAT (6X, 'TOTAL COMPOSITION:', T40, F8.4)
190
с
       CALCULATE THE MOLE FRACTION OF COMPONENT IN THE FEED STREAM
с
c
      DO I = 1, J
N(I)=M(I)/SUM
       END DO
C
       CHECK WHETHER THE FEED IS AT BUBBLE POINT OR DEW POINT.
с
ċ
       IF (FEED .EQ. 'DEW' .OR. FEED .EQ. 'dew') THEN
       GO TO 30
       ELSEIF (FEED .EQ. 'BUBBLE' .OR. FEED .EQ. 'bubble') THEN
       CALL BUBBLE
       GO TO 999
       ENDIF
       CALL DEW
30
      GO TO 999
      CLOSE (3,STATUS='KEEP')
CLOSE (1)
999
       STOP
       END
с
0000
       ***********
      THIS PROGRAM CALCULATES THE DEW POINT
       SUBROUTINE DEW
       COMMON/DATA/BP, DP, J, K, N
       REAL N(20), K(20), DP(20) , BP(20)
       DPT = 0.0
       DO I = 1, J
DP(I)=N(I)/K(I)
       DPT=DPT+N(I)/K(I)
       END DO
      200
       DO I =1, J
```

```
WRITE (1, 210)I, N(I), DP(I)
FORMAT (12X, I2, 16X, F8.4, 20X, F8.4)
 210
        END DO
       WRITE (1, 220)DPT
FORMAT(1H, 76(1H-),/,6X,'TOTAL DEW POINT:', T59, F8.4)
 220
       IF (DPT .GE. 1.0) THEN GO TO 10
        ELSE
        WRITE (1, 230)
 230
        FORMAT(/,6X,'LOWER THE DEW-POINT TEMPERATURE: of')
        ENDIF
       GO TO 20
 10
       WRITE (1, 240)
FORMAT(/,6X,'RAISE THE DEW-POINT TEMPERATURE: of')
 240
с
       FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.
 20
       WRITE (1, *) CHAR(12)
       RETURN
        END
С
С
0000
        THIS PROGRAM CALCULATES THE BUBBLE-POINT
        SUBROUTINE BUBBLE
       COMMON/DATA/BP, DP, J, K, N
       REAL N(20), K(20), DP(20), BP(20)
       BPT=0.0
       DO I = 1, J
BP(I) =N(I)*K(I)
       BPT=BPT+N(I)*K(I)
       END DO
       250
       DO I =1 ,J
WRITE (1, 260) I, N(I), BP(I)
FORMAT (12X, I2, 16X, F8.4, 20X, F8.4)
 260
       WRITE (1, 270)BPT
FORMAT(1H, 76(1H-),/,6X,'TOTAL BUBBLE POINT:', T59, F8.4)
270
       IF (BPT .GE. 1.0) THEN GO TO 30
       ELSE
       WRITE (1, 280)
280
       FORMAT(/,6X, 'RAISE THE BUBBLE POINT TEMPERATURE: oF')
```

	ENDIF GO TO 35
30 290	WRITE (1, 290) FORMAT(/,6X,'LOWER THE BUBBLE POINT TEMPERATURE: of')
с	FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.
35	WRITE (1, *) CHAR(12)
	RETURN END

PROGRAM PROG72

00000000000	EQUILIBRIUM FLASH CALCULATIONS THE PROGRAM USES THE ITERATIVE CONVERGENCE METHOD SUGGESTED BY I.H. OLIVER AND LATER MODIFIED BY S.T. KOSTECKE TO CALCULATE FLASH VAPORIZATION CONTAINING UP TO 15 COMPONENTS OF MULTICOMPONENT MIXTURES OF OIL, GAS AND CHEMICAL PROCESSES. THIS PROGRAM FURTHER CHECKS THE FEED COMPOSITION AT THE CONDITIONS GIVEN TO DETERMINE WHETHER THE FEED IS ALL VAPOR OR ALL LIQUID. THIS PROGRAM IS BASED ON THE WELL-KNOWN AND WIDELY USED VAPORIZATION EQUILIBRIUM RELATIONSHIPS THAT APPEAR IN HANDBOOKS BY THE NATURAL GAS PROCESSORS SUPPILIERS ASSOCIATIONS AND IN OTHER LITERATURE.
с с с с с	MI = TOTAL MOLES OF COMPONENT I IN THE FEED STREAM KI = EQUILIBRIUM FLASH CONSTANT FOR EACH COMPONENT I IN THE FEED STREAM R = L/V RATIO
	R1 = NEW L/V RATIO FOR EACH ITERATIVE CALCULATION L = TOTAL MOLES OF LIQUID AT EQUILIBRIUM CONDITIONS V = TOTAL MOLES OF VAPOR AT EQUILIBRIUM CONDITIONS Ni = MOLES FRACTION OF COMPONENT i IN THE FEED STREAM Li = TOTAL MOLES OF COMPONENT i IN THE LIQUID PHASE Vi = TOTAL MOLES OF COMPONENT i IN THE VAPOR PHASE X = MOLE FRACTION OF FEED IN THE LIQUID PHASE AT EQUILIBRIUM Y = MOLE FRACTION OF FEED IN THE VAPOR PHASE AT EQUILIBRIUM F, = CONSTANTS USED BY KOSTECKS
c	DIMENSION C(15), V(15), X1(15), Y1(15) REAL K(15), L1, L(15), M(15), N(15)
	COMMON/DATA1/ NUM, T, P, M, K COMMON/DATA2/ ADD1, ADD2, ADD3
	OPEN (UNIT = 3, FILE='DATA72.DAT', STATUS='OLD', ERR=18) OPEN (UNIT = 1, FILE='PRN')
с с с с с	READ FLASH TEMPERATURE, OF = T READ FLASH PRESSURE, psia = P READ NUMBER OF COMPONENTS = NUM READ FEED AND EQUILIBRIUM CONSTANT OF EACH COMPONENT = M, K
	READ (3, *, ERR=19) T READ (3, *, ERR=19) P READ (3, *, ERR=19) NUM
	DO I = 1, NUM READ (3, *, ERR=19) M(I), K(I) END DO
	GO TO 10
18 100	WRITE (*, 100) Format (5x, 'data file does not exist') Go to 999

```
WRITE (*, 110)
FORMAT (5X, 'ERROR MESSAGE IN THE DATA VALUE')
GO TO 999
19
110
         WRITE(1, 120)
FORMAT(//,20X,'MULTICOMPONENT EQUILIBRIUM FLASH CALCULATIONS')
 10
 120
         SUM=0.0
         DO I = 1, NUM
SUM = SUM + M(I)
         END DO
         CALCULATE MOLE FRACTION OF COMPONENT IN THE FEED STREAM
с
         DO I = 1, NUM
N(I)=M(I)/SUM
         END DO
         TOTAL=0.0
         DO I = 1, NUM
               TOTAL = TOTAL + N(I)
         END DO
         CHECK WHETHER THE FEED IS ALL LIQUID OR VAPOR
с
         SUM1=0.0
         DO I1 = 1, NUM
SUM1 = SUM1 + N(I1)*K(I1)
          END DO
         IF ( SUM1 .GT. 1.0) THEN
GO TO 45
          ELSE
               GO TO 900
         ENDIF
         SUM2 = 0.0
 45
         DO I2 = 1, NUM
SUM2 = SUM2+N(I2)/K(I2)
          END DO
          IF(SUM2 .GT. 1.0) THEN
GO TO 60
ELSE
          GO TO 1000
          ENDIF
         GUESS R=L/V
с
       R=1.0
SUM3=0.0
 60 .
65
```

DO I3 = 1, NUM C(I3) = (M(I3)*(K(I3)-1.0)*(R+1.0))/(K(I3)+R)END DO с CALCULATE E AND F CONSTANTS USED BY KOSTECKE. E = 0.0F = 0.0DO I4 = 1, NUM E = E+C(I4) F = F+(C(I4)**2/M(I4)) END DO с CALCULATE R1, THE NEW L/V RATIO R1=ABS((F*R-E*(R+1.0))/(F+E*(R+1.0))) IF (ABS(E) .LT. 0.001) THEN GO TO 90 ELSE R=R1 GO TO 65 ENDIF 90 L1 = 0.0V1 = 0.0Do I5 = 1, NUM L(I5) = (M(I5)*R1)/(K(I5)+R1) V(I5) = M(I5)-L(I5) L1 = L1 + L(I5) V1 = V1 + V(I5) END DO R2=L1/V1 CALCULATE X: MOLE FRACTION OF FEED IN THE LIQUID PHASE CCCC AT EQUILIBRIUM, AND Y: MOLE FRACTION OF FEED IN THE VAPOR PHASE AT EQUILIBRIUM. VAL1=0.0 VAL2=0.0 VAL3=0.0 DO I6 = 1, NUM VAL1 = VAL1+L(I6)VAL2 = VAL2+M(I6)VAL3 = VAL3 + V(16)END DO CALCULATE THE MOLE FRACTIONS IN THE LIQUID AND VAPOR PHASES. С С DO I = 1, NUM X1(I) =L(I)/VAL1 Y1(I) =V(I)/VAL3 END DO

C C C	CALCULATE THE TOTAL MOLE FRACTIONS OF THE FEED, LIQUID AND VAPOR PHASES
C	ADD1 = 0.0 ADD2 = 0.0 ADD3 = 0.0
с	DO I = 1, NUM ADD1 = ADD1+X1(I) ADD2 = ADD2+N(I) ADD3 = ADD3+Y1(I) END DO
c c	PRINT THE RESULTS ON THE SCREEN
C	CALL OUTPUT(N, L, X1, V, Y1, VAL1, VAL2, VAL3)
с	PRINT THE RESULTS ONTO THE PRINTER
	CALL PAPER (N, L, X1, V, Y1, VAL1, VAL2, VAL3)
	GO TO 999
900 300	WRITE(*,300) FORMAT(3X,'THE FEED COMPOSITION IS ALL LIQUID') GO TO 999
1000 400	WRITE(*,400) Format(3X,'THE FEED COMPOSITION IS ALL VAPOR')
	CLOSE (UNIT=3, STATUS='KEEP') CLOSE (UNIT=1)
999	STOP END
000000	**************************************
L.	SUBROUTINE OUTPUT (N, L, X1, V, Y1, VAL1, VAL2, VAL3) DIMENSION X1(15), Y1(15), V(15) REAL K(15), M(15), N(15), L(15)
	COMMON/DATA1/ NUM, T, P, M, K COMMON/DATA2/ ADD1, ADD2, ADD3
130	WRITE(*, 130) FORMAT(2X,'MULTICOMPONENT EQUILIBRIUM FLASH CALCULATION AT '\)
140	WRITE(*, 140) T, P FORMAT(F6.1,1X,'DEG. F',1X,'AND',1X,F6.1,1X,'FSIA')
150	WRITE(*, 150) FORMAT(70(1H*))

```
150 FORMAT(79(1H*))
```

160	WRITE(*, 160) FORMAT(/,1X,'COMPONENT',1X,'K-VALUE',6X,'FEED',16X,'LIQUID ',\)
170	WRITE(*, 170) Format(15X,'VAPOR')
180	WRITE(*, 180) FORMAT(1X,'NUMBER',12X,'Mols/h.',2X,'Mol. frac.',2X,'Mols/h.',\)
190	WRITE(*, 190) FORMAT(2X,'Mol.frac.',2X,'Mols/h.',2X,'Mol.frac.')
200	WRITE(*, 200) Format(78(1H-))
210	DO I = 1, NUM WRITE(*, 210) I, K(I), M(I), N(I), L(I), X1(I), V(I) FORMAT(1X,I2,5X,F5.3,2X,F9.3,2X,F8.3,3X,F9.3,2X,F8.3,2X,F9.3\)
220	WRITE(*, 220) Y1(I) FORMAT(2X,F8.3) END DO
230	WRITE(*, 230) Format(78(1H-))
240	WRITE(*, 240) VAL2, ADD2, VAL1, ADD1 Format(2X,'TOTALS',7X,F9.3,5X,F5.3,3X,F9.3,5X,F5.3,\)
250	WRITE(*, 250) VAL3, ADD3 Format(2X,F9.3,5X,F5.3)
260	WRITE(*, 260) Format (78(1H-)) Return END
с с с с	**************************************
	SUBROUTINE PAPER (N, L, X1, V, Y1, VAL1, VAL2, VAL3)
	DIMENSION V(15), X1(15), Y1(15) REAL K(15), L(15), M(15), N(15)
	COMMON/DATA1/NUM, T, P, M, K COMMON/DATA2/ADD1, ADD2, ADD3
270	WRITE (1, 270) FORMAT (/,2X,'MULTICOMPONENT EQUILIBRIUM FLASH CALCULATION AT ', $)$
280	WRITE (1, 280) T, P FORMAT (P6.1, 1X, 'DEG. F', 1X, 'AND', 1X, F6.1, 1X, 'psia')
290	WRITE (1, 290) Format (80(1H*))

300	<pre>WRITE (1, 300) FORMAT (/, 1X,'COMPONENT', 1X,'K-VALUE',6X,'FEED',16X,'LIQUID ',\)</pre>
310	WRITE (1, 310) Format (15X, 'VAPOR')
320	<pre>WRITE (1, 320) FORMAT (1X, 'NUMBER',12X,'Mols/h.',2X,'Mol. frac.',2X,'Mols/h.',\)</pre>
330	WRITE (1, 330) FORMAT (2X,'Mol. frac.', 2X, 'Mols/h.', 2X,'Mol. frac.')
340	WRITE (1, 340) FORMAT (80(1H-))
	DO I = 1, NUM
350	WRITE (1, 350) I, K(I), M(I), N(I), L(I), X1(I), V(I) FORMAT (1X,I2,5X,F5.3,2X,F9.3,2X,F8.3,3X,F9.3,2X,F8.3,2X,F9.3,\)
360	WRITE (1, 360) Y1(I) Format (2X,F8.3)
	END DO
370	WRITE (1, 370) Format (80(1H-))
380	WRITE (1, 380) VAL2, ADD2, VAL1, ADD1 FORMAT (2X,'TOTALS', 7X, F9.3, 5X, F5.3, 3X, F9.3, 5X, F5.3,\)
390	WRITE (1, 390) VAL3, ADD3 Format (2X, F9.3, 5X, F5.3)
400	WRITE (1, 400) Format (80(1H-))
с	FORM FEED THE PRINTING PAPER ON TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR(12)
	RETURN END

PROGRAM PROG73

000000000000000000000000000000000000000	TOWER SIZING FOR VALVE TRAYS. THE PROGRAM EMPLOYS THE NUTTER METHOD FOR CALCULATING TOWER DIAMETER WHERE VALVE-TYPE TRAYS ARE USED. TOWER SIZING GRAPHS AND CURVES IN NUTTER'S FLOAT VALVE DESIGN HAVE BEEN FITTED TO A SERIES OF EQUATIONS AND ARE USED IN THIS PROGRAM. ************************************
c c	REAL K, M, LCFS COMMON PI COMMON/MULT1/ X, SF, RD, V, VRD COMMON/MULT2/ AB, ADC, AT, DIA, RDCA COMMON/MULT3/ WV, WL, DENL, DENV, FF, REST, TS, SURT COMMON/MULT4/ LCFS, VCFS
	OPEN (UNIT=3,FILE='DATA73.DAT',STATUS='OLD',ERR=18) OPEN (UNIT=1,FILE='PRN')
00000000	READ THE VAPOR FLOW RATE, WV: lb/hr. READ THE LIQUID FOW RATE, WL: lb/hr. READ THE LIQUID DENSITY, DENL: lb/ft^3. READ THE VAPOR DENSITY, DENL: lb/ft^3. READ THE FOAM FACTOR, FF: READ THE DOWNCOMER RESIDENCE TIME, REST: sec. READ THE TRAY SPACING, TS: READ THE SURFACE TENSTION, SURT: dyne/cm.
	READ(3, *, ERR=19) WV, WL, DENL READ(3, *, ERR=19) DENV, FF, REST READ(3, *, ERR=19) TS, SURT

GO TO 20

18 21	WRITE (*, 21) Format(3X,'Data file does not exist') Go to 999
19 22	WRITE (*, 22) Format(3x,'error message in the data value') Go to 999
с с с	CONSTANTS FOR VELOCITY AT ZERO LIQUID LOAD, ft/sec. (W.W. BLACKWELL, Chemical Process Design On A Programmable Calculator, Mc. Graw-Hill, 1984).
20	PI = 3.1415927 A = 0.22982 B = 0.44605 C = 0.03452 D = 0.00415 E = 0.00017 K = 0.91146 M = -0.03821
с	CALCULATE THE RATIO OF SURFACE TENSION TO THE VAPOR DENSITY, X
	X=SURT/DENV
с	CALCULATE THE SAFETY FACTOR.
	SF=K*(X)**M
с	CALCULATE THE DENSITY RADICAL.
~	RD=SQRT(DENV/(DENL-DENV))
c c	V IS VALID FOR 0.1 < X < 3000.
с	CALCULATE THE VELOCITY AT ZERO LIQUID LOAD, ft/sec.
	VAL1=A+B*ALOG(X)-C*(ALOG(X))**2+D*(ALOG(X))**3-E*(ALOG(X))**4 V=EXP(VAL1) VRD=V*RD
	IF (VRD .GE. 0.5) THEN GO TO 23 ELSE ENDIF GO TO 24
23	VRD = 0.5
с	CHECK THE TRAY SPACING, TS.
24	IF (TS .EQ. 12.0) THEN
	GO TO 33
	ELSEIF (TS .EQ. 18.0) THEN

556

GO TO 44 ELSEIF (TS .EQ. 24.0) THEN GO TO 55 ELSEIF (TS .EQ. 30.0) THEN GO TO 66 ELSEIF (TS .EQ. 36.0) THEN GO TO 77 ENDIF 33 CALL PROG1 с PRINT THE RESULTS ON THE SCREEN CALL OUTPUT PRINT THE RESULTS ONTO A PRINTER. С CALL PAPER GO TO 999 44 CALL PROG2 с PRINT THE RESULTS ON THE SCREEN CALL OUTPUT С PRINT THE RESULTS ONTO A PRINTER CALL PAPER GO TO 999 55 CALL PROG3 с PRINT THE RESULTS ON THE SCREEN CALL OUTPUT С PRINT THE RESULTS ONTO A PRINTER CALL PAPER GO TO 999 66 CALL PROG4 PRINT THE RESULTS ON THE SCREEN с

C PRINT THE RESULTS ON THE SCRE

с PRINT THE RESULTS ONTO A PRINTER. CALL PAPER GO TO 999 77 CALL PROG5 с PRINT THE RESULTS ON THE SCREEN. CALL OUTPUT PRINT THE RESULTS ONTO A PRINTER. С CALL PAPER CLOSE (3, STATUS='KEEP') CLOSE (1) 999 STOP END с c c c TOWER SIZING FOR VALVE TRAYS USING A 12 INCH TRAY SPACING. ***************** ***** ** **** SUBROUTINE PROG1 REAL LCFS COMMON PI COMMON/MULT1/ X, SF, RD, V, VRD COMMON/MULT2/ AB, ADC, AT, DIA, RDCA COMMON/MULT3/ WV, WL, DENL, DENV, FF, REST, TS, SURT COMMON/MULT4/ LCFS, VCFS G = 0.77174H = -0.02964с CALCULATE THE TRAY SPACING FACTOR, TSF TSF = G+H*ALOG(X)с CALCULATE THE VAPOR VOLUMETRIC RATE, VCFS VCFS = WV/(DENV*3600.0)с CALCULATE THE LIQUID VOLUMETRIC RATE, LCFS LCFS = WL/(DENL*3600.0) IF (VRD .GT. 0.5) THEN VRD = 0.5 ELSE ENDIF с CALCULATE THE BUBBLING AREA, AB

AB = (VCFS*RD+LCFS)/(TSF*VRD*FF)

ADC = (REST*LCFS*12.0)/(FF*TS)С CALCULATE THE TOWER AREA, AT AT = (2.0 * ADC + AB) / SFCALCULATE THE TOWER DIAMETER, DIA. С DIA = SQRT((4.0*AT)/PI)с с CALCULATE THE DOWNCOMER AREA RATIO, RDCA с RDCA = ADC/(2.*ADC + AB)RETURN END C C TOWER SIZING FOR VALVE TRAYS USING AN 18 INCH TRAY SPACING. С SUBROUTINE PROG2 REAL LCFS COMMON PI COMMON/MULT1/ X, SF, RD, V, VRD COMMON/MULT2/ AB, ADC, AT, DIA, RDCA COMMON/MULT3/ WV, WL, DENL, DENV, FF, REST, TS, SURT COMMON/MULT4/ LCFS, VCFS G = 0.93655H = -0.02310с CALCULATE THE TRAY SPACING FACTOR, TSF TSF = G+H*ALOG(X)CALCULATE THE VAPOR VOLUMETRIC RATE, VCFS С VCFS = WV/(DENV*3600.0)С CALCULATE THE LIQUID VOLUMETRIC RATE, LCFS LCFS = WL/(DENL*3600.0)IF (VRD .GT. 0.5) THEN VRD = 0.5 ELSE ENDIF С CALCULATE THE BUBBLING AREA, AB. AB = (VCFS*RD+LCFS)/(TSF*VRD*FF) с CALCULATE THE DOWNCOMER AREA, ADC ADC = (REST*LCFS*12.0)/(FF*TS)

ADC = (REST*LCFS*12.0)/(FF*TS)CALCULATE THE TOWER AREA, AT. С AT = (2.0*ADC+AB)/SFс CALCULATE THE TOWER DIAMETER, DIA. DIA = SQRT((4.0*AT)/PI)с CALCULATE THE DOWNCOMER AREA RATIO, RDCA RDCA = ADC/(2.0*ADC+AB)RETURN END С c c TOWER SIZING FOR VALVE TRAYS USING A 24 INCH TRAY SPACING. SUBROUTINE PROG3 REAL LCFS COMMON PI COMMON/MULT1/ X, SF, RD, V, VRD COMMON/MULT2/ AB, ADC, AT, DIA, RDCA COMMON/MULT3/ WV, WL, DENL, DENV, FF, REST, TS, SURT COMMON/MULT3/ LCFS, VCFS c c CALCULATE THE TRAY SPACING FACTOR, TSF FOR ALL VALUES OF X, TSF=1.0 TSF = 1.0с CALCULATE THE VAPOR VOLUMETRIC RATE, VCFS VCFS = WV/(DENV*3600.0)с CALCULATE THE LIQUID VOLUMETRIC RATE, LCFS LCFS = WL/(DENL*3600.0)IF (VRD .GT. 0.5) THEN VRD = 0.5 ELSE ENDIF С CALCULATE THE BUBBLING AREA, AB AB = (VCFS*RD+LCFS)/(TSF*VRD*FF) с CALCULATE THE DOWNCOMER AREA, ADC ADC = (REST*LCFS*12.0)/(FF*TS)с CALCULATE THE TOWER AREA, AT.

AT = (2.0 * ADC + AB)/SF

```
CALCULATE THE TOWER DIAMETER, DIA.
С
       DIA = SQRT((4.0*AT)/PI)
С
        CALCULATE THE DOWNCOMER AREA RATIO, RDCA
       RDCA = ADC/((2.0*ADC)+AB)
        RETURN
        END
0000
        TOWER SIZING FOR VALVE TRAYS USING A 30 INCH TRAY SPACING.
        SUBROUTINE PROG4
        REAL LCFS
       COMMON PI
       COMMON/MULT1/ X, SF, RD, V, VRD
COMMON/MULT2/ AB, ADC, AT, DIA, RDCA
COMMON/MULT3/ WV, WL, DENL, DENV, FF, REST, TS, SURT
COMMON/MULT4/ LCFS, VCFS
        G = 0.98067
       H = 0.03220
с
       CALCULATE THE TRAY SPACING FACTOR, TSF
       TSF = G + H \star ALOG(X)
        IF ( X .LT. 1.85 ) THEN
TSF = 1.0
        ELSE
        ENDIF
       CALCULATE THE VAPOR VOLUMETRIC RATE, VCFS
с
        VCFS = WV/(DENV*3600.0)
с
        CALCULATE THE LIQUID VOLUMETRIC RATE, LCFS
        LCFS = WL/(DENL*3600.0)
        IF ( VRD .GT. 0.5) THEN
        VRD = 0.5
        ELSE
        ENDIF
с
       CALCULATE THE BUBBLING AREA, AB
        AB = (VCFS*RD+LCFS)/(TSF*VRD*FF)
с
        CALCULATE THE DOWNCOMER AREA, ADC.
        ADC =(REST*LCFS*12.0)/(FF*TS)
```

C CALCULATE THE TOWER AREA, AT

AT = (2.0*ADC+AB)/SF

C CALCULATE THE TOWER DIAMETER, DIA.

DIA = SQRT((4.0*AT)/PI)

C CALCULATE THE DOWNCOMER AREA RATIO, RDCA

RDCA = ADC/((2.0*ADC)+AB)

RETURN END

SUBROUTINE PROG5

```
REAL LCFS
COMMON PI
```

```
COMMON/MULT1/ X, SF, RD, V, VRD
COMMON/MULT2/ AB, ADC, AT, DIA, RDCA
COMMON/MULT3/ WV, WL, DENL, DENV, FF, REST, TS, SURT
COMMON/MULT4/ LCFS, VCFS
```

```
G = 0.96583
H = 0.06162
```

C CALCULATE THE TRAY SPACING FACTOR, TSF.

TSF = G+H*ALOG(X)IF (X .LT. 1.85) THEN

```
TSF = 1.0
ELSE
ENDIF
```

C CALCULATE THE VAPOR VOLUMETRIC RATE, VCFS

VCFS = WV/(DENV*3600.0)

C CALCULATE THE LIQUID VOLUMETRIC RATE, LCFS.

LCFS = WL/(DENL*3600.0)

```
IF (VRD .GT. 0.5) THEN
VRD = 0.5
ELSE
ENDIF
```

C CALCULATE THE BUBBLING AREA, AB.

AB = (VCFS*RD+LCFS)/(TSF*VRD*FF)

C CALCULATE THE DOWNCOMER AREA, ADC

ADC = (REST*LCFS*12.0)/(FF*TS)CALCULATE THE TOWER AREA, AT. C AT = (2.0*ADC+AB)/SFс CALCULATE THE TOWER DIAMETER, DIA. DIA = SQRT((4.0*AT)/PI)с CALCULATE THE DOWNCOMER AREA RATIO, RDCA RDCA = ADC/((2.0*ADC)+AB)RETURN END С с THIS PROGRAM PRINTS THE RESULTS ONTO THE SCREEN SUBROUTINE OUTPUT REAL LCFS COMMON/MULT1/ X, SF, RD, V, VRD COMMON/MULT2/ AB, ADC, AT, DIA, RDCA COMMON/MULT3/ WV, WL, DENL, DENV, FF, REST, TS, SURT COMMON/MULT4/ LCFS, VCFS WRITE (*, 100) FORMAT (//, 25X, 'TOWER DESIGN FOR VALVE TRAYS',/,78(1H*)) 100 WRITE (*, 110) WV, VCFS, WL, LCFS, DENL FORMAT (5X,'VAPOR FLOW RATE, lb/hr.:', T60, F10.1,/, 5X,'VAPOR VOLUMETRIC RATE, ft^3/sec.:', T60, F10.3,/, 5X,'LIQUID FLOW RATE, lb/hr.:', T60, F10.1,/, 5X,'LIQUID VOLUMETRIC RATE, ft^3/sec.:', T60, F10.3,/, 5X,'LIQUID DENSITY, lb/ft^3.:', T60, F10.3) 110 ٠ * WRITE (*, 120) DENV, FF, REST FORMAT (5X,'VAPOR DENSITY, 1b/ft^3.:', T60, F10.3,/, 5X,'FOAM FACTOR:', T60, F10.3,/, 5X,'DOWNCOMER RESIDENCE TIME, sec.:', T60, F10.3) 120 WRITE (*, 130) TS, SURT FORMAT (5X,'TRAY SPACING, in.:', T60, F8.0,/, 5X,'SURFACE TENSION, dyne/cm.:', T60, F8.2) 130 WRITE (*, 140) X, SF, RD FORMAT (5X,'SURFACE TENSION/VAPOR DENSITY (X):', T60, F10.3,/, 5X,'SAFETY FACTOR:', T60, F10.3,/, 5X,'DENSITY RADICAL:', T60, F10.3) 140 * WRITE (*, 150) AB, ADC, AT FORMAT (5X,'BUBBLING AREA, ft^2:', T60, F10.3,/, 5X,'DOWNCOMER AREA, ft^2:', T60, F10.3,/, 5X,'TOWER AREA, ft^2:', T60, F10.3) 150 WRITE (*, 160) DIA, RDCA FORMAT (5X,'TOWER DIAMETER, ft:', T60, F10.3,/, 160

	*	5X,'DOWNCOMER AREA RATIO:', T60, F10.4)
170		WRITE (*, 170) Format (78(1H-))
		RETURN END
0000		**************************************
с		SUBROUTINE PAPER
		REAL LCFS COMMON /MULT1/ X, SF, RD, V, VRD COMMON/ MULT2/ AB, ADC, AT, DIA, RDCA COMMON/ MULT3/ WV, WL, DENL, DENV, FF, REST, TS, SURT COMMON/ MULT4/ LCFS, VCFS
180		WRITE (1, 180) FORMAT (///, 25X, 'TOWER DESIGN FOR VALVE TRAYS',/,78(1H*))
190	* * * *	<pre>WRITE (1, 190) WV, VCFS, WL, LCFS, DENL FORMAT (5X,'VAPOR FLOW RATE, lb/hr.:', T60, F10.1,/, 5X,'VAPOR VOLUMETRIC RATE, ft'3/sec.:', T60, F10.3,/, 5X,'LIQUID FLOW RATE, lb/hr.:', T60, F10.1,/, 5X,'LIQUID VOLUMETRIC RATE, ft^3/sec.:',T60,F10.3,/, 5X,'LIQUID DENSITY, lb/ft^3.:', T60, F10.3)</pre>
200	*	WRITE (1, 200) DENV, FF, REST FORMAT (5X,'VAPOR DENSITY, lb/ft^3.:', T60, F10.3,/, 5X,'FOAM FACTOR:', T60, F10.3,/, 5X,'DOWNCOMER RESIDENCE TIME, sec.:', T60, F10.3)
210	*	WRITE (1, 210) TS, SURT FORMAT (5X,'TRAY SPACING, in.:', T60, F8.0,/, 5X,'SURFACE TENSION, dyne/cm.:', T60, F8.2)
220	*	WRITE (1, 220) X, SF, RD FORMAT (5X,'SURFACE TENSION/VAPOR DENSITY (X):', T60, F10.3,/, 5X,'SAFETY FACTOR:', T60, F10.3,/, 5X,'DENSITY RADICAL:', T60, F10.3)
230	* *	WRITE (1, 230) AB, ADC, AT FORMAT (5X,'BUBBLING AREA, ft^2:', T60, F10.3,/, 5X,'DOWNCOMER AREA, ft^2:', T60, F10.3,/, 5X,'TOWER AREA, ft^2:', T60, F10.3)
240	*	WRITE (1, 240) DIA, RDCA FORMAT (5X,'TOWER DIAMETER, ft:', T60, F10.3,/, 5X,'DOWNCOMER AREA RATIO:', T60, F10.4)
250		WRITE (1, 250) Format (78(1H-))
с		FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
		WRITE (1, *) CHAR(12)
		RETURN END

PROGRAM PROG74

с	PACKED-TOWER SIZING
č	THE TOWER SIZING METHOD IS BASED ON THE GENERALIZED PRESSURE
č	DROP CORRELATION OF SHERWOOD, SHIPLEY AND HOLLOWAY AND
с	REPLOTTED IN FIGURE GR-109 R-5 BY NORTON CO. THIS CORRELATION
с	IS A PLOT OF X .VS. Y FOR PRESSURE DROP CURVES RANGING FROM
с	0.5 TO 1.5 INCH OF WATER. FROM THESE CURVES THE TOWER DIAMETER
с	CAN BE EVALUATED.
с	***************************************
c c	L = LIOUID RATE. lb/h
c	L = LIQUID RATE, lb/h G = VAPOR RATE, lb/h
c	$DG = VAPOR DENSITY, 1b/ft^3$
č	$DL = LIQUID DENSITY, 1b/ft^3$
c	X = X CO-ORDINATE IN GR-109 R-5 PLOT
с	Y = Y CO-ORDINATE IN GR-109 R-5 PLOT
с	$G1 = VAPOR RATE, 1b/s.ft^2$
с	PF = PACKING FACTOR FOR PACKING USED
с	LVIS = LIQUID VISCOSITY, CP
с	$GC = GRAVITATIONAL CONSTANT = 32.2 It/s^2$
c c	PF = FACKING FACTOR FOR FOR FOR FACTOR FOR FOR FOR FACTOR FOR FOR FOR F
c	DELP = PRESSURE DROP inch H20/ft.
c	DELP = PRESSURE DROP inch H2O/ft. PI = CONSTANT = 3.1415927
č	*****
	COMMON/TOWER3/ AREA, DIA COMMON/TOWER5/ AREAF, DIA COMMON/TOWER5/ AREAF, DIAF REAL L, LVIS OPEN (UNIT = 3, FILE='DATA74.DAT', STATUS='OLD', ERR=18) OPEN (UNIT = 1, FILE='PRN')
000000	READ LIQUID FLOW RATE, lb/hr.=LREAD GAS FLOW RATE, lb/hr.=GREAD GAS DENSITY, lb/ft^3.=DGREAD LIQUID DENSITY, lb/ft^3.=DLREAD TOWER PACKING FACTOR=PFREAD TOWER PACKING FACTOR=DELPREAD LIQUID VISCOSITY, cP.=LVIS
	READ (3, *, ERR =19) L, G, DG READ (3, *, ERR =19) DL, PF, DELP, LVIS
	GO TO 10
18 100	WRITE (*, 100) Format (5x, 'data file does not exist') Go to 999
19	WRITE (*, 110)

110	FORMAT (5X, 'ERROR MESSAGE IN THE DATA VALUE') GO TO 999
10	PI = 3.1415927 GC = 32.2
с	WHERE 0.01 < X < 2.0 SUM1=DG/(DL-DG) X = L/G*SQRT(SUM1)
	IF (DELP .EQ. 0.05) THEN GO TO 30
	ELSEIF(DELP .EQ. 0.10) THEN GO TO 35
	ELSEIF(DELP .EQ. 0.25) THEN GO TO 40
	ELSEIF(DELP .EQ. 0.50) THEN GO TO 45
	ELSEIF(DELP .EQ. 1.0) THEN GO TO 50
	ELSEIF(DELP .EQ. 1.5) THEN GO TO 55
	ENDIF
30	CALL PROG1 (L, G, DG, DL, PF, LVIS)
с	PRINT THE RESULTS ON THE SCREEN
	CALL OUTPUT (L, G, DG, DL, PF, DELP, LVIS)
с	PRINT THE RESULTS ONTO A PRINTER
	CALL PAPER (L, G, DG, DL, PF, DELP, LVIS)
	GO TO 999
35	CALL PROG2 (L, G, DG, DL, PF, LVIS)
с	PRINT THE RESULTS ON THE SCREEN
	CALL OUTPUT (L, G, DG, DL, PF, DELP, LVIS)
с	PRINT THE RESULTS ONTO A PRINTER
	CALL PAPER (L, G, DG, DL, PF, DELP, LVIS) GO TO 999
40	CALL PROG3 (L, G, DG, DL, PF, LVIS)
с	PRINT THE RESULTS ON THE SCREEN

	CALL OUTPUT (L, G, DG, DL, PF, DELP, LVIS)
с	PRINT THE RESULTS ONTO A PRINTER
	CALL PAPER (L, G, DG, DL, PF, DELP, LVIS) Go To 999
45	CALL PROG4 (L, G, DG, DL, PF, LVIS)
с	PRINT THE RESULTS ON THE SCREEN
	CALL OUTPUT (L, G, DG, DL, PF, DELP, LVIS)
с	PRINT THE RESULTS ONTO A PRINTER
	CALL PAPER (L, G, DG, DL, PF, DELP, LVIS) Go To 999
50	CALL PROG5 (L, G, DG, DL, PF, LVIS)
с	PRINT THE RESULTS ON THE SCREEN
	CALL OUTPUT (L, G, DG, DL, PF, DELP, LVIS)
с	PRINT THE RESULTS ONTO A PRINTER
	CALL PAPER (L, G, DG, DL, PF, DELP, LVIS) GO TO 999
55	CALL PROG6 (L, G, DG, DL, PF, LVIS)
с	PRINT THE RESULTS ON THE SCREEN
	CALL OUTPUT (L, G, DG, DL, PF, DELP, LVIS)
с	PRINT THE RESULTS ONTO A PRINTER
	CALL PAPER (L, G, DG, DL, PF, DELP, LVIS) GO TO 999
	CLOSE (UNIT = 3, STATUS='KEEP') CLOSE (UNIT = 1)
999 C	STOP END
0000	**************************************
	SUBROUTINE PROG1 (L, G, DG, DL, PF, LVIS) REAL L,LVIS
	COMMON/TOWER1/ GC, PI COMMON/TOWER2/ X, Y, X1, Y1 COMMON/TOWER3/ AREA, DIA COMMON/TOWER4/ G1, G2, VG

COMMON/TOWER5/ AREAF, DIAF с CONSTANTS FOR DELP=0.05 C0 = -6.30253C1 = -0.60809C2 = -0.11932C3 = -0.00685C4 = 0.000320с WHERE 0.001 < Y < 0.15 SUM1 = ALOG(X)SUM2 = (ALOG(X)) * 2SUM3 = (ALOG(X))**3SUM4 = (ALOG(X))**4SUMT = C0+(C1*SUM1)+(C2*SUM2)+(C3*SUM3)+(C4*SUM4) Y1 = EXP(SUMT)CALCULATE VAPOR RATE 1b/s.ft^2 С VAL1 = Y1*DG*(DL-DG)*GC VAL2 = PF*(LVIS)**0.1RATIO = VAL1/VAL2G1 = (RATIO) * *0.5с CALCULATE TOWER AREA AND TOWER DIAMETER AREA = G/(3600.*G1)DIA = 12.0*((4.0*AREA)/PI)**0.5 CALCULATE Y CO-ORDINATE IN GR-109 R-5 PLOT С VAL3 = (G1**2)*PF*(LVIS)**0.1 VAL4 = DG*(DL+DG)*GC Y = VAL3/VAL4С CALCULATE THE PACKED TOWER FLOODING CONDITIONS CALL FLOOD (L, G, DG, DL, PF, LVIS) RETURN END с С PACKED-TOWER SIZING WITH A PRESSURE DROP OF 0.10 inch H2O/ft ***** с SUBROUTINE PROG2 (L, G, DG, DL, PF, LVIS) REAL L,LVIS COMMON/TOWER1/ GC, PI COMMON/TOWER2/ X, Y, X1, Y1 COMMON/TOWER3/ AREA, DIA COMMON/TOWER3/ G1, G2, VG COMMON/TOWER5/ AREAF, DIAF С CONSTANTS FOR DELP=0.1

	$\begin{array}{l} C0 &= -5.50093\\ C1 &= -0.78508\\ C2 &= -0.13496\\ C3 &= 0.00134\\ C4 &= 0.00174 \end{array}$
с	WHERE 0.001 < Y < 0.15
	SUM1 = ALOG(X) SUM2 = (ALOG(X))**2 SUM3 = (ALOG(X))**3 SUM4 = (ALOG(X))**4 SUMT = C0+(C1*SUM1)+(C2*SUM2)+(C3*SUM3)+(C4*SUM4) Y1 = EXP(SUMT)
с	CALCULATE VAPOR RATE, lb/s.ft^2
	VAL1 = Y1*DG*(DL-DG)*GC VAL2 = PF*(LVIS)**0.1 RATIO = VAL1/VAL2 G1 = (RATIO)**0.5
с	CALCULATE TOWER AREA AND TOWER DIAMETER
	AREA = G/(3600.*G1) DIA = 12.0*((4.0*AREA)/PI)**0.5
с	CALCULATE Y CO-ORDINATE IN GR-109 R-5 PLOT
	VAL3 = (G1**2)*PF*(LVIS)**0.1 VAL4 = DG*(DL-DG)*GC Y = VAL3/VAL4
с	CALCULATE THE PACKED TOWER FLOODING CONDITIONS.
	CALL FLOOD (L, G, DG, DL, PF, LVIS)
	RETURN END
с с с	**************************************
	SUBROUTINE PROG3 (L, G, DG, DL, PF, LVIS)
	REAL L, LVIS COMMON/TOWER1/ GC, PI COMMON/TOWER2/ X, Y, X1, Y1 COMMON/TOWER3/ AREA, DIA COMMON/TOWER4/ G1, G2, VG COMMON/TOWER5/ AREAF, DIAF
с	CONSTANTS FOR DELP=0.25
	C0 = -5.00319 C1 = -0.95299

 $C_2 = -0.13930$ $C_3 = 0.01264$

C4 = 0.00334WHERE 0.001 < Y < 0.15 с SUM1 = ALOG(X)SUM2 = (ALOG(X)) **2SUM3 = (ALOG(X)) **3SUM4 = (ALOG(X))**4 SUMT = C0+(C1*SUM1)+(C2*SUM2)+(C3*SUM3)+(C4*SUM4)Y1 = EXP(SUMT)С CALCULATE VAPOR RATE, 1b/s.ft^2 VAL1 = Y1*DG*(DL-DG)*GC VAL2 = PF*(LVIS)**0.1RATIO = VAL1/VAL2G1 = SQRT(RATIO)CALCULATE TOWER AREA AND TOWER DIAMETER с AREA = G/(3600.0*G1)DIA = 12.0*SQRT((4.0*AREA)/PI)с CALCULATE Y CO-ORDINATE IN GR-109 R-5 PLOT VAL3 = (G1**2)*PF*(LVIS)**0.1 VAL4 = DG*(DL-DG)*GC Y = VAL3/VAL4с CALCULATE THE PACKED TOWER FLOODING CONDITONS. CALL FLOOD (L, G, DG, DL, PF, LVIS) RETURN END с С PACKED TOWER SIZING WITH A PRESSURE DROP OF 0.5 inch H2O/ft с SUBROUTINE PROG4 (L, G, DG, DL, PF, LVIS) REAL L,LVIS COMMON/TOWER1/ GC, PI COMMON/TOWER2/ X, Y, X1, Y1 COMMON/TOWER3/ AREA, DIA COMMON/TOWER3/ AREA, DIA COMMON/TOWER5/ AREAF, DIAF с CONSTANTS FOR DELP=0.5 C0 = -4.39918 C1 = -0.99404 $\begin{array}{rcl} C1 & = -0.16983 \\ C2 & = -0.16983 \\ C3 & = & 0.00873 \\ C4 & = & 0.00343 \end{array}$ с WHERE 0.001 < Y < 0.15 SUM1 = ALOG(X)SUM2 = (ALOG(X)) **2

SUM3 = (ALOG(X))**3SUM4 = (ALOG(X))**4 SUMT = C0+(C1*SUM1)+(C2*SUM2)+(C3*SUM3)+(C4*SUM4) Y1 = EXP(SUMT)С CALCULATE VAPOR RATE, 1b/s.ft^2 VAL1 = Y1*DG*(DL-DG)*GC VAL2 = PF*(LVIS)**0.1 RATIO = VAL1/VAL2 G1 = (RATIO) **0.5С CALCULATE TOWER AREA AND TOWER DIAMETER AREA = G/(3600.*G1) DIA = 12.0*((4.0*AREA)/PI)**0.5 с CALCULATE Y CO-ORDINATE IN GR-109 R-5 PLOT VAL3 = (G1**2)*PF*(LVIS)**0.1 VAL4 = DG*(DL-DG)*GCY = VAL3/VAL4с CALCULATE THE PACKED TOWER FLOODING CONDITIONS. CALL FLOOD (L, G, DG, DL, PF, LVIS) RETURN END с c c PACKED-TOWER SIZING WITH A PRESSURE DROP OF 1 inch H2O/ft. ********************* *********************************** SUBROUTINE PROG5 (L, G, DG, DL, PF, LVIS) REAL L,LVIS COMMON/TOWER1/ GC, PI COMMON/TOWER2/ X, Y, X1, Y1 COMMON/TOWER3/ AREA, DIA COMMON/TOWER4/ G1, G2, VG COMMON/TOWER5/ AREAF, DIAF с CONSTANTS FOR DELP=1.0 C0 = -4.09505C1 = -1.00120 $\begin{array}{rcl} c_{2} &=& -0.15871\\ c_{3} &=& 0.00797\\ c_{4} &=& 0.00318 \end{array}$ с WHERE 0.001 < Y < 0.15 SUM1 = ALOG(X)SUM2 = (ALOG(X))**2SUM3 = (ALOG(X))**3SUM4 = (ALOG(X)) **4SUMT = C0+(C1*SUM1)+(C2*SUM2)+(C3*SUM3)+(C4*SUM4)Y1 = EXP(SUMT)

CALCULATE VAPOR RATE, 1b/s.ft^2 с VAL1 = Y1*DG*(DL-DG)*GC VAL2 = PF*(LVIS)**0.1 RATIO = VAL1/VAL2 G1 = (RATIO) * * 0.5С CALCULATE TOWER AREA AND TOWER DIAMETER AREA = G/(3600.*G1)DIA = 12.0*((4.0*AREA)/PI)**0.5 с CALCULATE Y CO-ORDINATE IN GR-109 R-5 PLOT VAL3 = (G1**2)*PF*(LVIS)**0.1 VAL4 = DG*(DL-DG)*GCY = VAL3/VAL4 с CALCULATE THE PACKED TOWER FLOODING CONDITIONS. CALL FLOOD (L, G, DG, DL, PF, LVIS) RETURN END с PACKED-TOWER SIZING WITH A PRESSURE DROP OF 1.5 inch H2O/ft. с с *************** SUBROUTINE PROG6 (L, G, DG, DL, PF, LVIS) REAL L, LVIS COMMON/TOWER1/ GC, PI COMMON/TOWER2/ X, Y, X1, Y1 COMMON/TOWER3/ AREA, DIA COMMON/TOWER3/ G1, G2, VG COMMON/TOWER5/ AREAF, DIAF с CONSTANTS FOR DELP=1.5 C0 = -4.02555C1 = -0.98945C2 = -0.08291 $C_3 = -0.03237$ $C_4 = 0.00532$ с WHERE 0.001 < Y < 0.15 SUM1 = ALOG(X)SUM2 = (ALOG(X)) **2SUM3 = (ALOG(X)) **3SUM4 = (ALOG(X)) **4SUMT = CO+(C1*SUM1)+(C2*SUM2)+(C3*SUM3)+(C4*SUM4)Y1 = EXP(SUMT)CALCULATE VAPOR RATE 1b/s.ft^2 с VAL1 = Y1*DG*(DL-DG)*GC VAL2 = PF*(LVIS)**0.1 RATIO = VAL1/VAL2

G1 = (RATIO) * *0.5

C CALCULATE TOWER AREA AND TOWER DIAMETER
AREA = G/(3600.*G1)
DIA = 12.0*((4.0*AREA)/PI)**0.5
C CALCULATE Y CO-ORDINATE IN GR-109 R-5 PLOT

C CALCULATE Y CO-ORDINATE IN GR-109 R-5 PLOT

VAL3 = (G1**2)*PF*(LVIS)**0.1 VAL4 = DG*(DL-DG)*GC Y = VAL3/VAL4

C CALCULATE THE PACKED TOWER FLOODING CONDITIONS.

CALL FLOOD (L, G, DG, DL, PF, LVIS) RETURN END

SUBROUTINE FLOOD (L, G, DG, DL, PF, LVIS)

REAL L,LVIS COMMON/TOWER// GC, PI COMMON/TOWER2/ X, Y, X1, Y1 COMMON/TOWER3/ AREA, DIA COMMON/TOWER4/ G1, G2, VG COMMON/TOWER5/ AREAF, DIAF

X1 = (L/G)*SQRT(DG/DL) TI = 62.4/DL Y1=10.0**(-1.6678-(1.085*ALOG10(X1))-(0.29655*(ALOG10(X1))**2)) VAL1 = Y1*DL*DG*GC VAL2 = PF*TT*(LVIS)**2 RATIO = VAL1/VAL2 G2 = SQRT(RATIO)

C CALCULATE VAPOR RATE AT 70% OF FLOODING CONDITION

G2 = 0.7*G2

C CALCULATE VAPOR VELOCITY AT 70% FLOODING CONDITION

VG=G2/DG

с

C CALCULATE THE TOWER AREA AT 70% FLOODING CONDITION.

C SINCE THE FLOODING CORRELATION IS NOT PERFECT, HENCE C TO HAVE A 95% CONFIDENCE, USE A SAFETY FACTOR OF 1.32 C FOR THE CALCULATED CROSS-SECTIONAL AREA (Bolles and Fair, C Chem. Eng., 89 (14), 109 (July 12, 1982).

AREAF = 1.32*(G/(3600.0*G2))

C CALCULATE THE TOWER DIAMETER AT 70% FLOODING CONDITION

DIAF = SQRT((4.0*AREAF)/PI)

с	DIAMETER IN inches:
	DIAF = 12.0*DIAF
	RETURN END
c c c	**************************************
	SUBROUTINE OUTPUT (L, G, DG, DL, PF, DELP, LVIS)
	REAL L,LVIS COMMON/TOWER2/ X, Y, X1, Y1 COMMON/TOWER3/ AREA, DIA COMMON/TOWER4/ G1, G2, VG COMMON/TOWER5/ AREAF, DIAF
120	WRITE (*, 120) Format(20X,'Packed Tower Design')
130	WRITE (*, 130) Format(78(1H*))
140	WRITE (*, 140) L Format(5x,'Llquid Flow Rate, lb/hr.:', T60, F12.3)
150	WRITE (*, 150) G Format(5x,'Gas flow rate, lb/hr.:', T60, F12.3)
160	WRITE (*, 160) DG Format(5X,'GAS DENSITY, lb/ft^3:', T60, F8.4)
170	WRITE (*, 170) DL FORMAT(5X,'LIQUID DENSITY, lb/ft^3:', T60, F8.4)
180	WRITE (*, 180) PF Format(5x,'Tower packing factor:', T60, F8.1)
190	WRITE (*, 190) DELP FORMAT(5X,'PRESSURE DROP, inch H2O/ft:', T60, F8.3)
200	WRITE (*, 200) LVIS Format(5x,'liquid Viscosity, cp:', T60, F8.3)
210	WRITE (*, 210) X FORMAT(5X,'X CO-ORDINATE IN THE GENERALIZED PLOT:', T60, F8.4)
220	WRITE (*, 220) Y FORMAT(5X,'Y CO-ORDINATE IN THE GENERALIZED PLOT:', T60, F8.4)
230	WRITE (*, 230) G1 FORMAT(5x,'CALCULATED VAPOR RATE, 1b/s.ft^2:', T60, F8.4)
240	WRITE (*, 240) X1 FORMAT(5X,'X CO-ORDINATE AT FLOODING CONDITION:', T60, F8.4)
	WRITE (*, 250) ¥1

250		FORMAT(5X,'Y CO-ORDINATE AT FLOODING CONDITION:', T60, F8.4)
260	*	WRITE (*, 260) G2 FORMAT(5X,'VAPOR RATE AT 70% FLOODING CONDITION, lb/s.ft^2:', T60, F8.4)
270	*	WRITE (*, 270) VG FORMAT(5X,'VAFOR VELOCITY AT 70% FLOODING CONDITION, ft./s.:', T60, F8.2)
280		WRITE (*, 280) AREA Format(5X,'Tower Area, ft^2:' T60, F8.2)
290		WRITE (*, 290) DIA FORMAT(5X,'TOWER DIAMETER, inches:', T60, F8.2)
300		WRITE (*, 300) AREAF FORMAT(5X,'TOWER AREA AT 70% FLOODING CONDITION, ft^2:',T60,F8.2)
310	*	<pre>WRITE (*, 310) DIAF FORMAT(5X,'TOWER DIAMETER AT 70% FLOODING CONDITION, inches:', T60, F8.2)</pre>
320 C		WRITE (*, 320) Format (78(1H-)) Return End
c		
с с		THIS PROGRAM PRINTS THE RESULTS ONTO A PRINTER

		SUBROUTINE PAPER (L, G, DG, DL, PF, DELP, LVIS)
		SUBROUTINE PAPER (L, G, DG, DL, PF, DELP, LVIS) REAL L, LVIS COMMON/TOWER2/ X, Y, X1, Y1 COMMON/TOWER3/ AREA, DIA COMMON/TOWER4/ G1, G2, VG
с		SUBROUTINE PAPER (L, G, DG, DL, PF, DELP, LVIS) REAL L, LVIS COMMON/TOWER2/ X, Y, X1, Y1 COMMON/TOWER3/ AREA, DIA COMMON/TOWER4/ G1, G2, VG COMMON/TOWER5/ AREAF, DIAF WRITE (1, 330)
C 330		SUBROUTINE PAPER (L, G, DG, DL, PF, DELP, LVIS) REAL L, LVIS COMMON/TOWER2/ X, Y, X1, Y1 COMMON/TOWER3/ AREA, DIA COMMON/TOWER4/ G1, G2, VG COMMON/TOWER5/ AREAF, DIAF WRITE (1, 330) FORMAT (///,25X,'PACKED TOWER DESIGN') WRITE (1, 340)
C 330 340		SUBROUTINE PAPER (L, G, DG, DL, PF, DELP, LVIS) REAL L, LVIS COMMON/TOWER2/ X, Y, X1, Y1 COMMON/TOWER3/ AREA, DIA COMMON/TOWER4/ G1, G2, VG COMMON/TOWER5/ AREAF, DIAF WRITE (1, 330) FORMAT (///,25X,'PACKED TOWER DESIGN') WRITE (1, 340) FORMAT (78(1H*)) WRITE (1, 350) L
C 330 340 350		SUBROUTINE PAPER (L, G, DG, DL, PF, DELP, LVIS) REAL L, LVIS COMMON/TOWER2/ X, Y, X1, Y1 COMMON/TOWER3/ AREA, DIA COMMON/TOWER4/ G1, G2, VG COMMON/TOWER5/ AREAF, DIAF WRITE (1, 330) FORMAT (///,25X,'PACKED TOWER DESIGN') WRITE (1, 340) FORMAT (78(1H*)) WRITE (1, 350) L FORMAT (5X,'LIQUID FLOW RATE, 1b/hr.:', T60, F12.3) WRITE (1, 360) G
C 330 340 350 360		<pre>SUBROUTINE PAPER (L, G, DG, DL, PF, DELP, LVIS) REAL L, LVIS COMMON/TOWER2/ X, Y, X1, Y1 COMMON/TOWER3/ AREA, DIA COMMON/TOWER5/ AREAF, DIA WRITE (1, 330) FORMAT (///,25X,'PACKED TOWER DESIGN') WRITE (1, 340) FORMAT (78(1H*)) WRITE (1, 350) L FORMAT (5X,'LIQUID FLOW RATE, lb/hr.:', T60, F12.3) WRITE (1, 360) G FORMAT (5X,'GAS FLOW RATE, lb/hr.:', T60, F12.3) WRITE (1, 370) DG</pre>

390	FORMAT (5X,'TOWER PACKING FACTOR:', T60, F8.1)
400	WRITE (1, 400) DELP Format (5X,'Pressure drop, INCH-H20:', T60, F8.3)
410	WRITE (1, 410) LVIS Format (5X,'Liquid Viscosity, cP:', T60, F8.3)
420	WRITE (1, 420) X FORMAT (5X,'X CO-ORDINATE IN THE GENERALIZED PLOT:', T60, F8.4)
430	WRITE (1, 430) Y Format (5x,'y co-ordinate in the generalized plot:', T60, F8.4)
440	WRITE (1, 440) G1 Format (5X,'Calculated Vapor Rate, 1b/s.ft^2:', T60, F8.4)
450	WRITE (1, 450) X1 FORMAT (5X,'X CO-ORDINATE AT FLOODING CONDITION:', T60, F8.4)
460	WRITE (1, 460) Y1 Format (5x,'y co-ordinate at flooding condition:', T60, F8.4)
470 *	WRITE (1, 470) G2 FORMAT (5X,'VAPOR RATE AT 70% FLOODING CONDITION, 1b/s.ft^2:', T60, F8.4)
480 *	<pre>WRITE (1, 480) VG FORMAT (5X,'VAPOR VELOCITY AT 70% FLOODING CONDITION, ft./sec:',</pre>
490	WRITE (1, 490) AREA Format (5X,'Tower Area, ft^2:', T60, F8.2)
500	WRITE (1, 500) DIA FORMAT (5X,'TOWER DIAMETER, inches:', T60, F8.2)
510 *	WRITE (1, 510) AREAF FORMAT (5X,'TOWER AREA AT 70% FLOODING CONDITION, ft^2:', T60,F8.2)
520 F *	WRITE (1, 520) DIAF ORMAT (5X,'TOWER DIAMETER AT 70% FLOODING CONDITION, inches:', T60, F8.2)
530	WRITE (1, 530) Format (78(1H-))
с	FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR(12) RETURN END

PROGRAM PROG75

с	NUMBER OF STAGES AT LOW CONCENTRATIONS.
с	THIS PROGRAM CALCULATES THE NUMBER OF EQUILIBRIUM STAGES
с	USING THE SMOKER EQUATIONS.
c	IN THE LOW-CONCENTRATION REGION THE EQUILIBRIUM RELATION FOLLOWS
c c	HENRY'S LAW AND Y=KX, AND THIS RELATIONSHIP IS USED TO PLOT THE EQUILIBRIUM LINE IF DATA ARE NOT AVAILABLE. THE EQUATIONS
c	DEVELOPED BY SMOKER WILL HANDLE THESE OPERATIONS ANALYTICALLY.
c	DEVILOPED DI SMOKAK WILL MANDLE IMEDI OF MAIITOND MAMETIONIZIO
č	*******
č	A1 = AVERAGE RELATIVE VOLATILITY OF THE RECTIFYING SECTION
C	A2 = AVERAGE RELATIVE VOLATILITY OF THE STRIPPING SECTION
С	R = REFLUX RATIO
с	XF = MOLE FRACTION OF THE FEED
с	XD = MOLE FRACTION OF THE TOP PRODUCT
с	XB = MOLE FRACTION OF THE BOTTOMS PRODUCT
с	NI = NUMBER OF STAGES IN THE RECTIFYING SECTION
c	N2 = NUMBER OF STAGES IN THE STRIPPING SECTION TOTAL = THE TOTAL NUMBER OF STAGES.
С	
c	POST = THE OPTIMUM FEED STAGE LOCATION.
c c	******
c	
C	REAL K, K1, K2, N1, N2
	COMMON/MASS/K1,K2
	OPEN (UNIT=3, FILE='DATA75.DAT', STATUS='OLD', ERR=18)
	OPEN (UNIT=1, FILE='PRN')
	••••••
с	READ THE RELATIVE VOLATILITIES OF THE TWO COMPONENTS A1, A2.
с	READ THE MOLE FRACTION OF THE FEED, XF.
с	READ THE MOLE FRACTION OF THE DISTILLATE, XD.
с	READ THE MOLE FRACTION OF THE BOTTOMS, XB.
с	READ THE REFLUX RATIO, R.
	READ (3, *, ERR=19) A1, A2
	READ (3, *, ERR=19) XF, XD, XB
	READ (3, *, ERR=19) R
	GO TO 5
18	WRITE (*, 23)
23	FORMAT (5X, 'DATA FILE DOES NOT EXIST')
20	GO TO 999
19	WRITE (*, 29)
29	FORMAT (5X, 'ERROR MESSAGE IN THE DATA VALUE')
	GO TO 999
5	WRITE (1, 100)
	FORMAT(//,15X,49HEQUILIBRIUM NUMBER OF STAGES BY SMOKER'S EQUATION,
	/1H ,78(1H*))
	/ /· //
	WRITE (1, 110) A1, A2
110	FORMAT(5X,'THE RELATIVE VOLATILITY OF COMPONENT 1.:', T60,
*	F10.3,/,5X,'THE RELATIVE VOLATILITY OF COMPONENT 2.:',

T60,F10.3)

*

-	160,710.3)
120 * *	<pre>WRITE (1, 120) XF, XD, XB, R FORMAT(5X,'THE MOLE FRACTION OF THE FEED:', T60, F10.4,/, 5X,'THE MOLE FRACTION OF THE DISTILLATE:', T60, F10.4,/, 5X,'THE MOLE FRACTION OF THE BOTTOMS:', T60, F10.4,/, 5X,'THE REFLUX RATIO:', T60, F10.2)</pre>
	GO TO 10
С	CALCULATE THE NUMBER OF STAGES IN THE RECTIFYING SECTION
10	S1=R/(R+1.0) B1=XD/(R+1.0)
C C C	CALCULATE K, THE VALUE OF THE X-ORDINATE AT THE POINT WHERE THE EXTENDED OPERATING LINE INTERSECT THE VAPOUR-LIQUID EQUILIBRIUM CURVE FROM THE QUADRATIC EQUATION $AX^2+BX+C=0$
c c	CALCULATE THE CO-EFFICIENTS OF THE QUADRATIC EQUATION IN THE RECTIFYING SECTION
	A = S1*(A1-1.0) B = S1+B1*(A1-1.0)-A1 C = B1
с	SOLVE THE QUADRATIC EQUATION
	CALL PROG1 (A, B, C, K)
с	CALCULATE THE NUMBER OF STAGES IN THE RECTIFYING SECTION, N1
	C1 = 1.0+(A1-1.0)*K X1 = XD-K X2 = XF-K BETA1 = (S1*C1*(A1-1.0))/(A1-(S1*(C1**2))) VAL1 = (X1*(1.0-(BETA1*X2)))/(X2*(1.0-(BETA1*X1))) VAL2 = A1/(S1*(C1**2))
с	CALCULATE THE NUMBER OF STAGES IN THE RECTIFYING SECTION.
	N1=ALOG(VAL1)/ALOG(VAL2)
130 *	WRITE (1, 130) N1 FORMAT(5X,'THE NUMBER OF STAGES IN THE RECTIFYING SECTION:', T60, F8.2)
с	CALCULATE THE NUMBER OF STAGES IN THE STRIPPING SECTION, N2
	S2 = ((R*XF)+XD-(R+1.0)*XB)/((R+1.0)*(XF-XB)) B2 = ((XF-XD)*XB)/((R+1.0)*(XF-XB))
c c	CALCULATE THE CO-EFFICIENTS OF THE QUADRATIC EQUATION AX^2+BX+C=0 IN THE STRIPPING SECTION

```
A = S2*(A2-1.0)
        B = (S2+B2*(A2-1.0)-A2)
        C = B2
        CALCULATE THE NUMBER OF STAGES IN THE RECTIFYING SECTION.
С
        CALL PROG1 (A, B, C, K)
        X3 = XF-K
        X4 = XB-K
        C2 = 1.0+(A2-1.0)*K
        BETA2 = (S2*C2*(A2-1.0))/(A2-(S2*(C2**2)))
        VAL3 = (X3*(1.0-(BETA2*X4)))/(X4*(1.0-(BETA2*X3)))
        VAL4 = \dot{A}2/(\dot{S}2*(C2**2))
       N2 = ALOG(VAL3)/ALOG(VAL4)
       WRITE (1, 140) N2
FORMAT(5X,'THE NUMBER OF STAGES IN THE STRIPPING SECTION:',
T60, F8.2)
140
       TOTAL = N1+N2
       WRITE (1, 150) TOTAL
FORMAT(5X,'TOTAL NUMBER OF STAGES:', T60, F8.2)
150
С
        THE FEED LOCATION IS THAT FROM THE STRIPPING SECTION TO
с
        THE BOTTOM OF THE COLUMN.
        POST = N2
       WRITE (1, 160) POST
FORMAT (5X,'FEED STAGE LOCATION:', T60, F8.1)
160
       WRITE (1, 170)
170
       FORMAT (78(1H-))
с
       FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.
       WRITE (1, *) CHAR(12)
        CLOSE (3,STATUS='KEEP')
        CLOSE (1)
999
        STOP
        END
С
        С
с
        THIS PROGRAM SOLVES THE QUADRATIC EQUATION AX^2+BX+C=0 IN THE
        RECTIFYING AND STRIPPING SECTIONS
с
        *******
с
                                               ***********************
       SUBROUTINE PROG1 (A, B, C, K)
       REAL K, K1, K2
       COMMON/MASS/K1,K2
       D = (B**2-4.0*A*C)
```

IF (D .LT. 0.0) THEN GO TO 10 ELSE K1= (-B+SQRT(D))/(2.0*A)K2= (-B-SQRT(D))/(2.0*A)ENDIÈ IF (K1 .LT. 0.0 .OR. K1 .GT. 1.0) THEN GO TO 20 ELSE K = K1GO TO 30 ENDIF IF (K2 .LT. 0.0 .OR. K2 .GT. 1.0) THEN GO TO 40 20 ELSE K=K2 GO TO 30 ENDIF WRITE (*, 190) FORMAT(6X,'ROOTS ARE OUT OF LIMITS CHECK PROBLEM SPECIFICATION') 40 190 GO TO 30 WRITE (*, 200) FORMAT(6X, 'IMAGINARY ROOTS, CHECK PROBLEM SPECIFICATION') 10 200 30 RETURN END

PROGRAM PROG76

с	RAPID E	STIM	ATES OF MULTICOMPONENT DISTILLATION AND
с	MINIMUM	TRA	YS IN DISTILLATION COLUMNS
с	THIS PRO	OGRAJ	M WILL CALCULATE THE FEED COMPONENT RECOVERY
с	IN BOTH	THE	COLUMN DISTILLATE AND BOTTOMS PRODUCTS AND
c	ESTIMAT	ETH	E MINIMUM NUMBER OF STAGES NEEDED TO PERFORM
č	THE GIVEN SEPARATION.		
č			S CONTAINING UP TO 12 COMPONENTS CAN BE HANDLED
c	BY THIS		
	BI THIS	PRO	JRAM.
С			
с	THE FOL	LOWI	NG ASSUMPTIONS FOR THIS PROGRAM ARE:
с			
с	1. THE	RE I	S ONLY ONE FEED STREAM WITH 12 OR FEWER COMPONENTS.
с	2. THE	RE I	S ONLY ONE HEAVY KEY COMPONENT.
с	3. FEE	D CO	MPONENTS ARE ARRANGED IN ORDER OF DECREASING
с	REL	ATIV	E VOLATILITY, WITH THE LIGHT KEY (LK) AND HEAVY KEY
с			MPONENTS SHOWN ADJACENT TO EACH OTHER.
č			D ENTERS THE COLUMN AT THE OPTIMAL STAGE.
č			UMN PRODUCES TWO PRODUCTS (DISTILLATE AND BOTTOMS)
c			ERHEAD CONDENSER AND BOTTOM REBOILER.
c			**************************************
c	ALPHAI		
c		=	EQUILIBRIUM CONSTANT FOR COMPONENT I IN THE FEED STREAM
с	кнк	-	
с			IN THE FEED STREAM
с	Di	-	COMPONENT i IN DISTILLATE, MOL
с		=	COMPONENT I IN BOTTOMS, MOL
с	BHK	=	HEAVY KEY COMPONENT IN BOTTOMS, MOL
с	FHK	=	HEAVY KEY COMPONENT IN THE FEED, MOL
с	DLK	=	
c	FLK	=	LIGHT KEY COMPONENT IN THE FEED, MOL
č	ALPHA (LI		
č	Fi		COMPONENT I IN THE FEED
c	NUM1		MINIMUM TRAYS AT TOTAL REFLUX
c		=	
č	BLK		LIGHT KEY COMPONENT IN BOTTOMS, MOL
с		=	
с	Q	=	FEED THERMAL CONDITION (RATIO OF LIQUID TO VAPOR
с			FOR THE FEED)
с	RECLK		THE PERCENTAGE RECOVERY OF THE LIGHT KEY COMPONENT
с	RECHK		THE PERCENTAGE RECOVERY OF THE HEAVY KEY COMPONENT
с	FACTOR		FACTOR FOR THE REFLUX RATIO (i.e. FACTOR*MREF)
с		=	THE POSITION OF THE FEED PLATE
с	CHOICE	-	DETERMINE WHETHER FEED IS EQUILIBRIUM CONSTANT OR
С			RELATIVE VOLATILITY.
с	******	****	**************
с			
	CHARACT	ER*5	CHOICE
	DIMENSI	ON A	LPHA(20), B(20), D(20), F(20), PERD(20), PERB(20)
			F(20), XB(20), XD(20)

DIMENSION XF(20), XB(20), XD(20) REAL K(20), MB(20), MD(20), NUM1, MREF

 $\tt COMMON/DATA1/NUM, F, K, RECLK, RECHK, K1, Q, FACTOR COMMON/DATA2/THETA, FEED$

OPEN (UNIT = 3, FILE='DATA76.DAT', STATUS='OLD', ERR=18) OPEN (UNIT =1, FILE='PRN')

```
с
000000000
        READ THE CHOICE OF PARAMETER e.g. RELATIVE VOLATILITY (REL) OR
        EQUILIBRIUM CONSTANT (EQUIL).
        READ THE TOTAL NUMBER OF COMPONENTS, NUM
        READ THE FEED AND RELATIVE VOLATILITY OR EQUILIBRIUM CONSTANT, F,K
        READ THE RECOVERIES OF LIGHT AND HEAVY KEYS, RECLK, RECHK.
READ THE POSITION OF THE HEAVY KEY COMPONENT, K1.
        READ THE THERMAL FEED CONDITION, Q
        READ THE FACTOR FOR THE REFLUX RATIO, FACTOR.
        READ (3, 5, ERR=19) CHOICE
 5
        FORMAT (A)
        IF (CHOICE .EQ. 'REL' .OR. CHOICE .EQ. 'rel') THEN
с
        READ THE TOTAL NUMBER OF COMPONENTS
        READ (3, *, ERR=19) NUM
С
        READ THE FEED AND THE RELATIVE VOLATILITY OF EACH COMPONENT.
        DO I = 1, NUM
        READ (3, *, ERR=19) F(I), K(I)
        END DO
        ELSEIF (CHOICE .EQ. 'EQUIL' .OR. CHOICE .EQ. 'equil') THEN
        READ THE TOTAL NUMBER OF COMPONENTS.
с
        READ (3, *, ERR=19) NUM
с
        READ THE FEED AND THE EOUILIBRIUM CONSTANT OF EACH COMPONENT.
        DO I = 1, NUM
        READ (3, *, ERR=19) F(I), K(I)
        END DO
        ENDIF
с
        READ THE RECOVERIES OF THE LIGHT AND HEAVY COMPONENTS.
        READ (3, *, ERR=19) RECLK, RECHK
        READ THE POSITION OF THE HEAVY KEY COMPONENT.
С
        READ THE THERMAL CONDITION OF THE FEED.
С
c
        READ THE FACTOR FOR THE REFLUX RATIO.
        READ (3, *, ERR=19) K1, Q, FACTOR
        GO TO 10
        WRITE (*, 22)
FORMAT (5X, 'DATA FILE DOES NOT EXIST')
 18
 22
        GO TO 999
```

19 23	WRITE (*, 23) Format (5x, 'error message in the data value') Go to 999
с с	ARRANGE THE FEEDS AND EQUILIBRIUM CONSTANT/RELATIVE VOLATILITY IN THE ORDER OF DECREASING VALUE
10	CALL SORT (K, F, NUM)
c c	CHECK WHETHER EQUILIBRIUM CONSTANTS OR RELATIVE VOLATILITY IS CHOSEN WITH THE FEED COMPONENTS
	IF (CHOICE .EQ. 'EQUIL' .OR. CHOICE .EQ. 'equil') THEN GO TO 15 ELSEIF (CHOICE .EQ. 'REL' .OR. CHOICE .EQ. 'rel') THEN
	DO I = 1, NUM ALPHA(I)=K(I) END DO ENDIF GO TO 25
c c	CALCULATE THE RELATIVE VOLATILITIES OF THE COMPONENTS EQUILIBRIUM VALUES OF THE LIGHT KEY AND HEAVY KEY COMPONENTS ARE:
15	DO I = 1, NUM ALPHA(I)=K(I)/K(K1) END DO
с	CALCULATE THE CORRELATION CONSTANTS: A1,B1
25	A1=ALOG10((1.0-RECHK)/RECHK) VAL1=(RECLK/(1.0-RECLK))*(RECHK/(1.0-RECHK)) B1=ALOG10(VAL1)/ALOG10(ALPHA(K1-1))
с	CALCULATE THE COMPONENT MOLES IN THE DISTILLATE
	DO I = 1, NUM D(I)=((10**A1)*(ALPHA(I)**B1))/(1.0+((10**A1)*(ALPHA(I)**B1))) PERD(I)=D(I)*100.0 MD(I)=D(I)*F(I) END DO
c c	CALCULATE THE RECOVERY OF THE LIGHT KEY COMPONENT IN THE DISTILLATE
	K2=K1-1 D(K2)=F(K2)*RECLK
с	CALCULATE THE TOTAL MOLES IN THE DISTILLATE
	SUM1=0.0
	DO I = 1, NUM

SUM1=SUM1+MD(I) END DO с CALCULATE THE PERCENTAGE RECOVERY IN THE BOTTOMS PRODUCT DO I = 1, NUM B(I)=1.0/(1.0+((10**A1)*(ALPHA(I)**B1))) PERB(I)=B(I)*100.0 $MB(I) = \hat{B}(I) * \hat{F}(I)$ END DO С CALCULATE THE RECOVERY OF THE HEAVY KEY COMPONENT IN THE BOTTOMS B(K1)=F(K1)*RECHK с CALCULATE THE TOTAL PRODUCT IN THE BOTTOMS SUM2=0.0 DO I = 1, NUM SUM2=SUM2+MB(I) END DO с CALCULATE THE MINIMUM NUMBER OF TRAYS IN THE COLUMN VAL2=(MD(K1-1)/(MD(K1))*(MB(K1)/MB(K1-1))) AMIN=ALOG(VAL2)/(ALOG(ALPHA(K1-1))) RECLK=RECLK*100.0 RECHK=RECHK*100.0 с CALCULATE THE MOLE FRACTIONS IN THE FEED, DISTILLATE AND BOTTOMS SUMB=0.0 SUMD=0.0 SUMF=0.0 DO J = 1, NUM SUMB=SUMB+MB(J) SUMD=SUMD+MD(J) SUMF=SUMF+F(J) END DO DO I = 1, NUM XB(I)=MB(I)/SUMB XD(I)=MD(I)/SUMD XF(I)=F(I)/SUMF END DO CALL UNDERW (NUM, Q, XF, XD, THETA, MREF, ALPHA, K1)

C USE MCCORMICK'S EQUATION TO DETERMINE GILLILAND'S GRAPHICAL C CORRELATION AND HENCE THE EQUILIBRIUM NUMBER OF STAGES.

```
REF=FACTOR*MREF
       X=(REF-MREF)/(REF+1.0)
C=0.105*ALOG10(X)+0.44
       Y=1.0-(X**C)
       NUM1=(Y+AMIN)/(1.0-Y)
c
c
       USE KIRKBRIDE'S EMPIRICAL EQUATION TO ESTIMATE THE POSITION
       OF THE FEED PLATE.
       Z1=(XF(K1)/XF(K1-1)*(SUM2/SUM1)*(XB(K1-1)/XD(K1))**2)
       C1=0.206*ALOG10(Z1)
       Z2=10**C1
       FEED=(Z2*NUM1)/(1.0+Z2)
с
      CALL OUTPUT(NUM, F, ALPHA, PERD, MD, PERB, MB, K1, RECLK, RECHK, AMIN, SUN1, SUM2, MREF, REF, NUM1)
    *
       CLOSE (3, STATUS='KEEP')
       CLOSE (1)
 999
       STOP
       END
С
       С
       THIS PROGRAM CALCULATES THE FEED COMPONENTS IN ORDER
c
c
       OF DECREASING RELATIVE VOLATILITY.
       **********************************
       SUBROUTINE SORT (X, Y, N)
       DIMENSION X(20), Y(20)
           N1=N-1
       DO 10 I≠1,N1
           I1≂I+İ
       DO 20 J=I1,N
          IF(X(I) .GE. X(J)) THEN
       GO TO 20
       ELSE
       HOLD=X(I)
       X(I)=X(J)
X(J)=HOLD
          HOLD=Y(I)
           Y(I)=Y(J)
           Y(J)=HOLD
           ENDIF
20
       CONTINUE
10
       CONTINUE
       RETURN
       END
С
       THIS PROGRAM CALCULATES THE UNDERWOOD CONSTANT THETA, USING THE
с
       BINARY SEARCH METHOD AND THE MINIMUM REFLUX RATIO
(i.e. AT AN INFINITE NUMBER OF PLATES)
с
с
с
```

```
SUBROUTINE UNDERW (N, Q, XF, XD, THETA, MREF, ALPHA, K1) DIMENSION ALPHA(20), XF(20), XD(20)
        REAL MREF, LHS
        THETA1=ALPHA(K1)
        THETA2=ALPHA(K1-1)
        DO 10 I=1,20
        THETA=0.5*(THETA1+THETA2)
        LHS=Q-1.0
        DO J = 1, N
        LHS=LHS+((ALPHA(J)*XF(J))/(ALPHA(J)+THETA))
        END DO
        IF (LHS .LT. 0.0) THEN
        THETA1=THETA
        GO TO 10
        ELSEIF (LHS .GT. 0.0) THEN
        THETA2=THETA
        GO TO 10
ELSEIF (LHS .EQ. 0.0) THEN
        GO TO 50
        ENDIF
 10
        CONTINUE
 50
        MREF=-1.0
        DO K = 1, N
MREF=MREF+((ALPHA(K)*XD(K))/(ALPHA(K)-THETA))
        END DO
        RETURN
        END
С
      C
C
      THIS PROGRAM OUTPUTS THE RESULTS ON THE SCREEN
      c
      SUBROUTINE OUTPUT(NUM, F, ALPHA, PERD, MD, PERB, MB, K1, RECLK, RECHK, AMIN,
     *SUM1, SUM2, MREF, REF, NUM1)
        DIMENSION F(20), ALPHA(20), PERD(20), PERB(20)
REAL MD(20), MB(20), NUM1, MREF
COMMON/DATA2/THETA, FEED
        WRITE (*, 100)
        FORMAT(25X, 'MULTICOMPONENT SYSTEM FRACTIONATION')
100
        WRITE (*, 110)
        FORMAT(80(1H*))
110
        WRITE (*, 120)
FORMAT(/,1X,'COMPONENT',7X,'FEED',5X,'REL VOLATILITY',3X,
'DISTILATTE ',8X,'BOTTOMS ')
120
       WRITE (*, 130)
FORMAT(1X,'NUMBER',9X,'MOLES',7X,'ALPHA',10X,'%',8X,'MOLES',8X,
130
     *'%',6X,'MOLES')
```

140	WRITE (*, 140) Format (80(1H-))
	DO 10 I=1,NUM
150 10	WRITE (*, 150) I,F(I),ALPHA(I),PERD(I),MD(I),PERB(I),MB(I) FORMAT(3X,I2,7X,F9.4,3X,F9.4,3X,F9.4,2X,F9.4,3X,F9.4,2X,F9.4) CONTINUE
160	WRITE (*, 160) Format (80(1H-))
170	WRITE (*, 170) K1 Format(3x,'The heavy key component number is:',T60,I2)
180 *	WRITE (*, 180) RECLK FORMAT(3X,'PERCENTAGE RECOVERY OF THE LIGHT KEY COMPONENT',/, 3X,'IN THE DISTILLATE IS (%):',T60,F7.2)
190 *	WRITE (*, 190) RECHK FORMAT(3X,'PERCENTAGE RECOVERY OF THE HEAVY KEY COMPONENT',/, 3X,'IN THE BOTTOMS IS (%):',T60,F7.2)
200	WRITE (*, 200) AMIN FORMAT(3X,'MINIMUM NUMBER OF STAGES IS:',T60,F9.1)
210 *	WRITE (*, 210) SUM1, SUM2 FORMAT(3X,'TOTAL MOLES IN THE DISTILLATE IS:',T60,F9.4,/, 3X,'TOTAL MOLES IN THE BOTTOMS IS:',T60,F9.4)
220	WRITE (*, 220) THETA FORMAT(3X,'UNDERWOOD CONSTANT:',T60,F9.4)
230	WRITE (*, 230) MREF Format(3X,'Minimum Reflux Ratio:',T60,F9.4)
240	WRITE (*, 240) REF Format(3x,'actual reflux ratio:',t60,f9.4)
250	WRITE (*, 250) NUM1 FORMAT(3X,'NUMBER OF THEORETICAL PLATES IN THE COLUMN:',T60,F9.1)
260	WRITE (*, 260) FEED Format(3x,'The position of the feed plate is:',T60,F4.1)
270	WRITE (*, 270) Format (1H ,80(1H-))
	OPEN (UNIT=1, FILE='LPT1')
*	CALL PAPER (NUM, F, ALPHA, PERD, ND, PERB, NB, K1, RECLK, RECHK, AMIN, SUM1, SUM1, SUM2, MREF, REF, NUM1)
	CLOSE (1) RETURN END
c c	THIS PROGRAM PRINTS THE RESULTS ONTO A PRINTER

с	************
*	SUBROUTINE PAPER(NUM,F,ALPHA,PERD,MD,PERB,MB,K1,RECLK,RECHK, AMIN,SUM1,SUM2,MREF,REF,NUM1)
	DIMENSION F(20), ALPHA(20), PERD(20), PERB(20) Real MD(20), MB(20), MREF, NUN1
	COMMON/DATA2/THETA, FEED
280 *	WRITE (1, 280) FORMAT (///,25X,'MULTICOMPONENT SYSTEM FRACTIONATION',/,1H , 80(1H*))
290 *	WRITE (1, 290) FORMAT (/, 1X,'COMPONENT',7X,'FEED',5X,'REL VOLATILITY', 3X, 'DISTILLATE ', 8X, 'BOTTOMS ')
300 *	WRITE (1, 300) FORMAT (1X,'NUMBER',9X,'MOLES',7X,'ALPHA',10X,'%',8X,'MOLES',8X, '%',6X,'MOLES')
310	WRITE (1, 310) Format (80(1H-))
	DO 10 I = 1 ,NUM
320	WRITE (1, 320) I, F(I), ALPHA(I), PERD(I), MD(I), PERB(I), MB(I) FORMAT (3X,I2,7X,F9.4,3X,F9.4,3X,F9.4,2X,F9.4,3X,F9.4,2X,F9.4)
10	CONTINUE
330	WRITE (1, 330) Format (80(1H-))
340	WRITE (1, 340) K1 Format (3X,'The heavy key component number:', T60, I2)
350 *	WRITE (1, 350) RECLK FORMAT (3X,'PERCENTAGE RECOVERY OF THE LIGHT KEY COMPONENT',/, 3X,'IN THE DISTILLATE (%):', T60, F7.2)
360 *	WRITE (1, 360) RECHK FORMAT (3X,'PERCENTAGE RECOVERY OF THE HEAVY KEY COMPONENT',/, 3X,'IN THE BOTTOMS (%):', T60, F7.2)
370	WRITE (1, 370) AMIN Format (3x,'Minimum number of stages:', T60, F9.1)
380 *	WRITE (1, 380) SUM1, SUM2 FORMAT (3X,'TOTAL MOLES IN THE DISTILLATE:', T60, F9.4,/, 3X,'TOTAL MOLES IN THE BOTTOMS:', T60, F9.4)
390	WRITE (1, 390) THETA FORMAT (3X,'UNDERWOOD CONSTANT:', T60, F9.4)
400	WRITE (1, 400) MREF Format (3x,'Minimum Reflux Ratio:', T60, F9.4)
	WRITE (1, 410) REF

.

410	FORMAT (3X,'ACTUAL REFLUX RATIO:', T60, F9.4)
420 *	WRITE (1, 420) NUM1 FORMAT (3X,'NUMBER OF THEORETICAL PLATES IN THE COLUMN:', T60, F9.1)
430	WRITE (1, 430) FEED FORMAT (3X,'THE POSITION OF THE FEED PLATE:', T60, F9.1)
440	WRITE (1, 440) Format (80(1H-))
с	FORM FEED THE PRINTING PAPER TO THE TOP OF NEXT PAGE.
	WRITE (1, *) CHAR(12) RETURN

•

END

CHAPTER 8

Heat Transfer

INTRODUCTION

The increasing cost of energy in recent years has resulted in increased attention to conservation and efficient energy management. Other types of technology, for example, pinch technology, have been employed in the energy integration of process plants and of heat exchangers in particular. This has often resulted in an improved performance of the plants at reduced operational costs. Various types of forced convective heat transfer equipment are used in the chemical process industries. These range from the simple shell and tube exchangers, double pipe, finned tube and tubular air cooled exchangers to the plate types, spiral and compact heat exchangers. Table 8-1 lists the major types of heat exchanger [1].

BASIC THEORY

Heat is transferred from points of high temperature to points of lower temperature by direct contact of particles of matter or the emission and absorption of radiant energy. The three classifications of heat transfer are:

Conduction. The transfer of heat from one body to another by physical contact. The rate of heat flow by conduction is proportional to the area available for the heat transfer and the temperature gradient in the direction of the heat flow path. The rate of heat flow in a given direction can be expressed as:

$$\frac{\mathrm{dQ}}{\mathrm{d\theta}} = -\mathrm{kA} \,\frac{\mathrm{dT}}{\mathrm{dx}} \tag{8-1}$$

Table 8-1	
Summary of Types of Heat Exchangers	Summary

Туре	Major Characteristics	Application
Shell and tube	Bundle of tubes encased in a cylindrical shell.	Always the first type exchanger to consider.
Air cooled heat exchanger	Rectangular tube bundles mounted on frame, with air used as the cooling medium	Economic where cost of cooling water is high.
Double pipe	Pipe within a pipe; inner pipe may be finned or plain.	For small units.
Extended surface	Externally finned tube.	Services where the outside tube resistance is appreciably greater than the inside resistance. Also used in debottlenecking existing units
Brazed plate fin	Series of plates separated by corrugated fins.	Cryogenic services: all fluids must be clean.
Spiral wound	Spirally wound tube coils within a shell	Cryogenic services: fluids must be clean.
Scrapped surface	Pipe within a pipe, with rotating blades scraping the inside walls of the inner pipe.	Crystallization cooling applications.
Bayonet tube	Tube element consists of an outer and inner tube.	Useful for high temperature difference between shell and tube fluids.
Falling film coolers	Vertical units using a thin film of water in tubes.	Special cooling applications.
Barometric condenser	Direct contact of water and vapor.	Where mutual solubilities of water and process fluid permit
Cascade coolers	Cooling water flows over series of tubes.	Special cooling applications for very corrosive process fluids.
Impervious graphite	Constructed of graphite for corrosion protection.	Used in very highly corrosive heat exchange services.

Source: Cheremisinoff [1]

where Q = amount of heat transferred in time θ hr, Btu

- k = the thermal conductivity, or proportionality constant, Btu/(hr)(ft²)(°F/ft)
- A = area of heat transfer perpendicular to direction of heat flow, ft^2
- T = temperature, $^{\circ}F$
- x =length of conduction path in direction of heat flow, ft

Convection. The transfer of heat by the mixing or movements of fluids or fluids with a solid. Mixing may occur as a result of density difference alone, as in natural convection. Alternatively, mechanically induced agitation may produce forced convection, as in turbulent flow in a heat exchanger tube, or to the heat transfer fluid in the jacket of an agitated vessel. The rate of heat transfer is:

$$\frac{\mathrm{d}Q}{\mathrm{d}\theta} = \mathrm{h}A\Delta t \tag{8-2}$$

where h = the heat transfer coefficient or the proportionality constant, which is a function of the type of agitation, the degree of turbulence, and the nature of the fluid. For the steady state condition, Equation 8-2 becomes

$$q = hA\Delta t \tag{8-3}$$

Radiation. The transfer of heat by the absorption of radiant energy, transformed into heat in surrounding bodies. The rate at which radiant heat energy is emitted from a source is:

$$\frac{\mathrm{d}Q}{\mathrm{d}\theta} = \sigma \varepsilon \mathrm{AT}^4 \tag{8-4}$$

where σ = Stefan-Boltzmann constant = 0.1713 × 10⁻⁸ Btu/hrft²(°R)⁴ ε = emissivity of surface A = exposed surface area of heat transfer, ft² T = absolute temperature, °R

The most common applications of heat transfer are a combination of one or more of these classifications. Also in many processes where a phase change occurs, that is, condensation, boiling, and evaporation, other physical phenomena play an important role in limiting the achievable heat transfer rate. However, only the most common, forced convective heat transfer, is considered here as well as an illustration of conductive heat transfer.

The procedures for the design of heat exchangers vary with the type of problem, but are often repetitive and tedious. An increasing number of software packages have been developed, for example, Heat Transfer and Fluid Flow Services (HTFS) and Heat Transfer Research Inc. (HTRI). These programs are available to the member companies. Two procedures are used in the design of heat exchangers. They are the rating method and the optimization design method.

The Rating Method

In the rating method, the designer assumes the form of an exchanger and then develops computations to determine whether the exchanger could handle the process requirements under known conditions. If the exchanger does not satisfy the requirements, a different exchanger is assumed, and the calculations repeated until a satisfactory design is obtained. For an example, for any given set of process requirements, the designer could assume an exchanger constructed of specific materials and having a known tube size, number of tubes, tube spacing, and passes, a shell size, baffling and number of passes (other more detailed assumptions such as removable bundle and floating head arrangements and end constructions). In addition, there are six process variables:

- inlet temperature of hot stream
- outlet temperature of hot stream
- inlet temperature of cold stream
- outlet temperature of hot stream
- flow rate of hot stream
- flow rate of cold stream

The designer will proceed by checking whether the energy balance calculations between the fluids are consistent. If there is a large deviation (for example 10%), the hot fluid duty is used for the design calculations. Further calculations could be the required heat transfer area, lengths, pressure drops on either side (ΔP), and an overall heat transfer coefficient. Repeated trials may be required to achieve an accurate overall heat transfer coefficient. If the results of the final design show that (allowing for any accumulations of scale during service between cleaning) the assumed exchanger is of reasonable dimensions, of a low cost, and will create acceptable pressure drops, the equipment is considered adequate for the duty.

Optimization Design Method

This approach is based upon optimum economic conditions. By choosing various conditions, the designer can arrive at a final design giving the least total cost for fixed charges and operation. Any increase in the fluid velocity on either side may result in a larger heat transfer coefficient, and consequently require a smaller heat transfer area and exchanger cost for a given rate of heat transfer. However, increased fluid velocity will result in a greater pressure drop and pumping costs. The optimum economic design occurs at the conditions where the total cost is at a minimum. This method minimizes the sum of the variable annual costs for the exchanger and its operation.

In general, the design of a heat exchanger involves the following steps:

- 1. Determine the flow rates and the rate of heat transfer required for given conditions.
- 2. Decide on the type of heat exchanger (e.g., 1:1, 1:2 type, or 2:4 type) to be used. Assume permissible scale factors and show the basic equipment specifications.
- 3. Calculate the fluid velocities of the film coefficients and the overall heat transfer coefficient.
- 4. Calculate the mean temperature difference driving force.
- 5. Determine the required area of heat transfer and the exchanger dimensions.
- 6. Evaluate the results to see that all dimensions, costs, ΔPs , and other design parameters are satisfactory.
- 7. If the results in Step 6 show that the exchanger is not satisfactory, the specifications given in Step 2 are inadequate. Choose new specifications and repeat Steps 3 through 7 until a satisfactory design is achieved.

This chapter presents rating calculations for the shell and tube heat exchanger, procedures for the design of double pipe or longitudinal finned tube heat exchanger, air cooler, heat tracer requirements for pipelines, and heat transfer in batch reactors.

LMTD IN SHELL AND TUBE HEAT EXCHANGERS

In a shell and tube heat exchanger, the rate of heat transfer from the hot to the cold fluid is directly proportional to the temperature difference between the two fluids. For the rating of heat exchangers, it is important to determine the mean temperature difference based on the hot and cold fluids' inlet and outlet temperatures. The heat transfer process is generally more efficient where there is countercurrent flow compared to co-current flow. The relative direction of the fluids affects the value of the log mean temperature difference (LMTD). A further advantage of approaching countercurrent flow, except where one fluid is isothermal (that is, constant temperature) is that in co-current flow, the hot fluid cannot be cooled below the cold fluid outlet temperature, so the ability to recover heat is limited. There are a few cases where co-current flow is preferred. For example, when cooling a viscous fluid, a higher heat transfer coefficient may be obtained by a rapid initial reduction in viscosity. Co-current may also be favorable if the temperature of the warmer fluid could otherwise be reduced to its freezing point.

In the majority of industrial operations, higher velocities, shorter tubes, and a more economical exchanger can be achieved with a multipass flow design as shown in Figure 8-1. The flow pattern is however part countercurrent and part co-current. Therefore, the mean temperature difference lies between the countercurrent and co-current log mean temperature differences (LMTD). The mean temperature differences for the

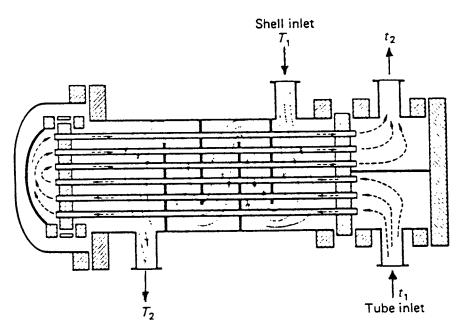


Figure 8-1. Flow patterns in a multipass exchanger (1:2 floating head as shown).

various types of crossflow exchanger are also less than for countercurrent flow designs, but greater than for co-current flow. In addition, the relationships between terminal temperatures and mean temperature difference restrict the performance of each type of exchanger. A correction factor, F, is defined to convert the LMTD to the corrected mean temperature difference (CMTD). For true countercurrent flow, F = 1, and as more co-current flow is introduced, the factor is reduced and the efficiency of the exchanger drops. A low value of F requires a large surface requirement for a given heat load. Performance can be enhanced by using several shells in series or by increasing the number of passes in the same shell. The lower limit of practical efficiency is in the range F = 0.75 to 0.8. The lowest F value is for one shell when rating shell and tube exchangers in series. This value is raised as the number of shells increases, and the flow more nearly resembles countercurrent flow. For the design of shell and tube exchangers, the minimum number of shells that will raise the F value above the chosen minimum of 0.75 to 0.8 is calculated.

Here, a computer program is written by setting the minimum correction factor to 0.75. The program automatically increases the number of shells until the value of F exceeds 0.75 or some other preset value.

Bowman *et al.* [2] simplified the calculations required to determine the mean temperature difference in a shell and tube exchanger. Figure 8-2 shows some of the major types of the shell and tube heat exchanger construction, and Figures 8-3 to 8-6 illustrate correction factor plots for various shell and tube exchangers. The equations given by Bowman *et al.* [2] are used to determine the corrected LMTD.

Heat Exchanger Design

The thermal rating of heat exchanger equipment requires the computation of the LMTD between the shell and tube fluids' temperatures. This is defined by

$$LMTD = \frac{\Delta t_1 - \Delta t_2}{\ln\left(\frac{\Delta t_1}{\Delta t_2}\right)}$$
(8-5)

where for countercurrent flow

 Δt_1 = the larger terminal temperature difference, $T_1 - t_2$

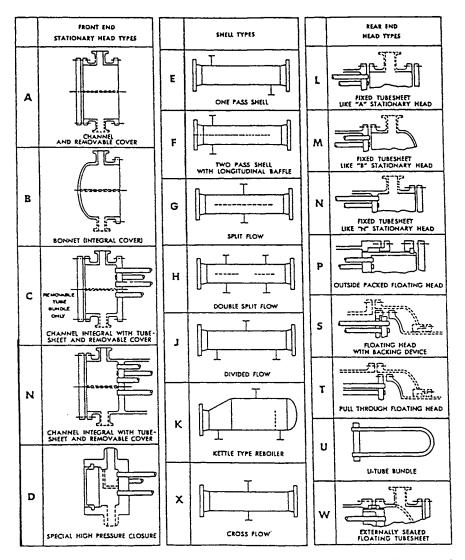


Figure 8-2. Tubular Exchanger Manufacturers Association (TEMA) classification and terminology for heat exchangers. TEMA heat exchanger head types.

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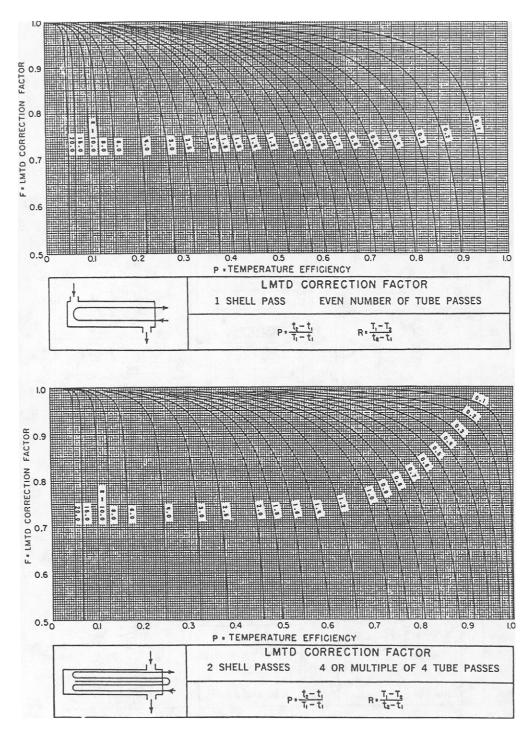


Figure 8-3. LMTD correction factor © 1978 Tubular Exchanger Manufacturers Association.

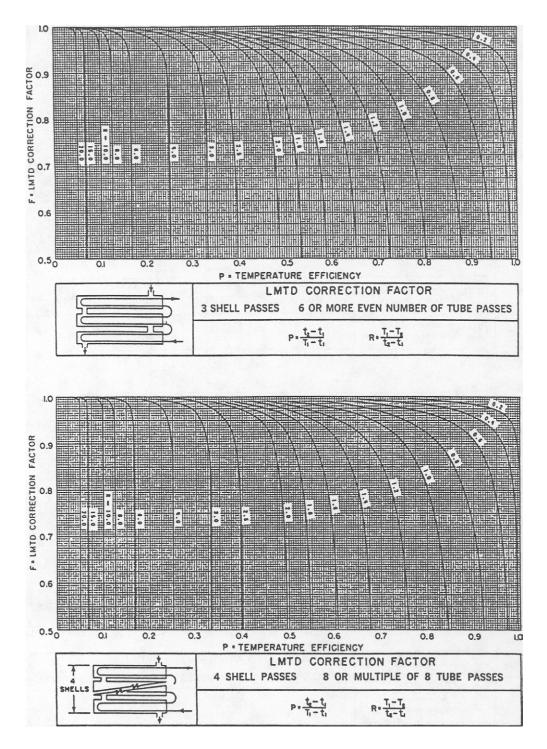


Figure 8-4. LMTD correction factor.

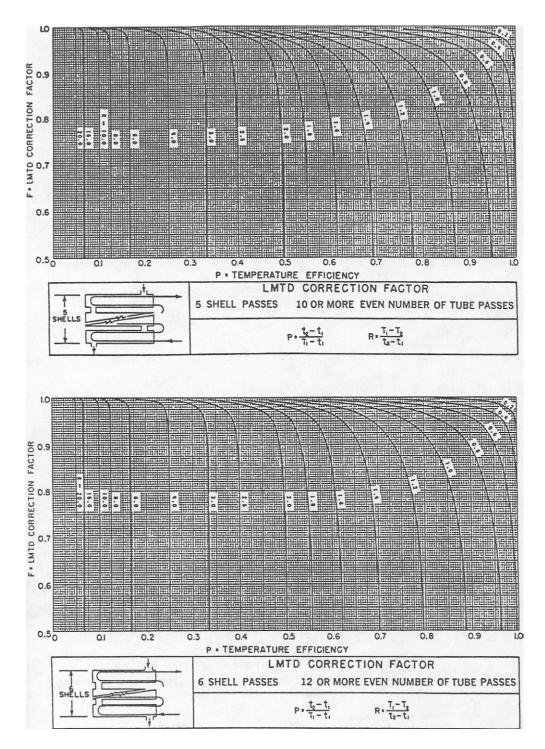
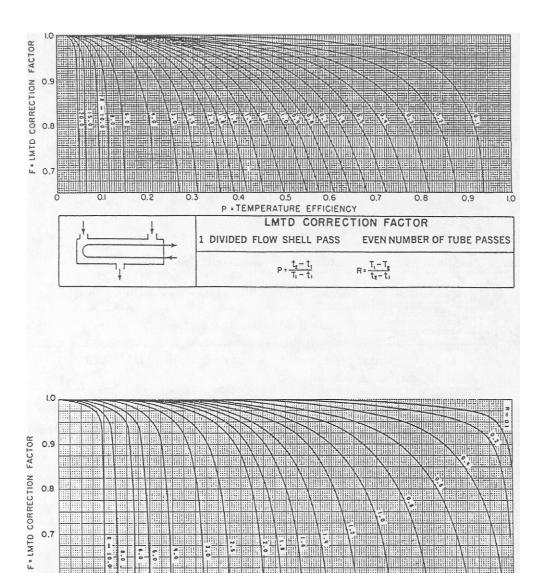


Figure 8-5. LMTD correction factor.



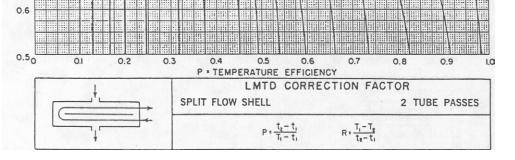


Figure 8-6. LMTD correction factor.

(text continued from page 596)

and

 Δt_2 = the smaller terminal temperature difference, $T_2 - t_1$

For co-current flow

 $\Delta t_1 = T_1 - t_1$

and

 $\Delta t_2 = T_2 - t_2$

The equations used to calculate F are based on the following assumptions:

- The overall heat transfer coefficient, U, is constant through the heat exchanger.
- The rate of flow of each fluid is constant.
- The specific heat of each fluid is constant.
- There is no condensation of vapor or boiling of liquid in that part of the exchanger considered.
- Heat losses are negligible.
- There is equal heat transfer surface in each pass.
- The temperature of the shell side fluid in any shell side pass is uniform over any cross section.

Figure 8-7 shows typical temperature profiles of two fluids in true countercurrent flow through a 1-1 exchanger.

$$P = \frac{t_2 - t_1}{T_1 - t_1}$$
(8-6)

$$R = \frac{T_1 - T_2}{t_2 - t_1}$$
(8-7)

For an exchanger with N shell passes, Bowman *et al.* [2] developed a general solution for the correction factor,

$$F = \frac{\left(\frac{\sqrt{R^{2} + 1}}{R - 1}\right) \ln\left\{\frac{(1 - P_{x})}{(1 - RP_{x})}\right\}}{\ln\left\{\frac{(2/P_{x}) - 1 - R + \sqrt{R^{2} + 1}}{(2/P_{x}) - 1 - R - \sqrt{R^{2} + 1}}\right\}}$$
(8-8)

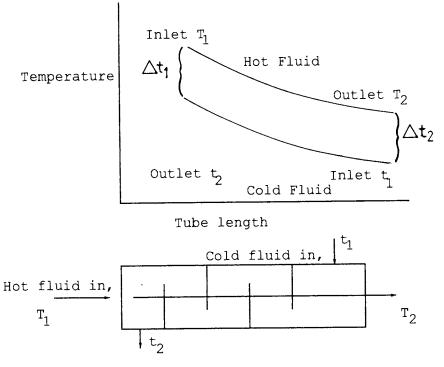


Figure 8-7. Counter-current flow in a 1-1 exchanger.

where

$$P_{x} = \frac{1 - \left[\frac{RP - 1}{P - 1}\right]^{1/N}}{R - \left[\frac{RP - 1}{P - 1}\right]^{1/N}}$$
(8-9)

N is the total number of shell passes, that is, the product of shell passes per shell and the number of units in series. Solving for N by repetitive trial and error with a minimum desired F, the minimum required number of shell passes can be determined.

If R = 1, Equation 8-9 becomes indeterminate, which reduces to

$$P_x = \frac{P}{(N - NP + P)}$$
(8-10)

Equation 8-8 then becomes

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$$F = \frac{\frac{P_x \sqrt{R^2 + 1}}{1 - P_x}}{\ln\left\{\frac{(2/P_x) - 1 - R + \sqrt{R^2 + 1}}{(2/P_x) - 1 - R - \sqrt{R^2 + 1}}\right\}}$$
(8-11)

The corrected mean temperature difference (CMTD) becomes

CMTD = (F)(LMTD)

(8-12)

Nomenclature

F = correction factor LMTD = log mean temperature difference, °F N = number of series exchanger shells required T_1 = hot fluid inlet temperature, °F T_2 = hot fluid outlet temperature, °F t_1 = cold fluid inlet temperature, °F t_2 = cold fluid outlet temperature, °F

Shell and Tube Heat Exchanger Design

The various designs of heat exchangers are well illustrated in standard texts and in the literature [3,4,5,6,7,8,9]. Recently, Pooley *et al.* [10] developed algorithms for both shell and tube exchangers and compact exchangers. Their algorithms rely on the allowable pressure drops of both of the streams being contacted.

The rating of an exchanger involves using the equations to calculate the suitability of an existing exchanger for given process conditions. The following steps are used for rating a shell and tube exchanger where there is no phase of change. The process conditions required are

Hot fluid: $T_1, T_2, W, C, s, \mu, k, R_d, \Delta P$

Cold fluid: $t_1, t_2, w, c, s, \mu, k, R_d, \Delta P$

The data required are

SHELL SIDE	TUBE SIDE
ID	Number and length
Baffle space	ID, OD, BWG, and pitch
passes	passes

From the process conditions, the heat load, Q, is

(1)
$$Q = WC(T_1 - T_2) = wc(t_2 - t_1)$$
 (8-13)

(2) The true temperature difference, Δt (assuming a number of tube passes) is:

LMTD =
$$\frac{(T_1 - t_2) - (T_2 - t_1)}{\ln\left\{\frac{T_1 - t_2}{T_2 - t_1}\right\}}$$
(8-14)

Approach factor, F_c , is determined from

$$\mathbf{P} = \frac{\mathbf{t}_2 - \mathbf{t}_1}{\mathbf{T}_1 - \mathbf{t}_1} \tag{8-6}$$

$$\mathbf{R} = \frac{\mathbf{T}_1 - \mathbf{T}_2}{\mathbf{t}_2 - \mathbf{t}_1} \tag{8-7}$$

and Figures 8-3 to 8-6. Alternatively, F_c can be calculated from Equation 8-8 and Equation 8-11.

$$(3) \Delta t = (LMTD)(F_c) \tag{8-15}$$

Tube Side

$$a_t = \frac{(no. of tubes)(flow area/tube)}{no. of pases}$$

$$= \frac{N_t \cdot a'_t}{144n}, \quad ft^2$$
 (8-16)

(5) Mass velocity, G_t, is

$$G_t = \frac{W}{a_t}, \quad \left(\frac{lb}{ft^2.hr}\right)$$
 (8-17)

(6) Reynolds number, Re_t , is

$$\operatorname{Re}_{t} = \frac{(\mathrm{ID}/12).\,\mathrm{G}_{t}}{\mu}$$
 (8-18)

⁽⁴⁾ Flow area a_t , ft^2 :

where

$$\mu = 2.42 \times cP \left(\frac{lb}{ft.hr.}\right)$$

(7) Prandtl number, Pr_t , is

$$\Pr_{t} = \frac{c\mu}{k} \tag{8-19}$$

(8) The heat transfer coefficient, h_i , for viscous flow (Re_t < 2100) as given by Sieder and Tate's correlation [11]:

$$\frac{\mathbf{h}_{i}.\,\mathrm{ID}}{\mathrm{k}} = 1.86 \left(\frac{\mathrm{ID}.\,\mathrm{G}}{\mu}\right)^{1/3} \left(\frac{\mathrm{c}\mu}{\mathrm{k}}\right)^{1/3} \left(\frac{\mathrm{ID}}{\mathrm{L}}\right)^{1/3} \left(\frac{\mu}{\mu_{\mathrm{w}}}\right)^{0.14}$$
(8-20)

where Nu = Nusselt number = h_i .ID/k ID = inside tube diameter, ft

Equation 8-20 can be expressed as:

$$\frac{\mathbf{h}_{i} \cdot \mathbf{ID}}{\mathbf{k}} = 1.86 \,\mathrm{Re}_{t}^{0.33} \,\mathrm{Pr}_{t}^{0.33} \left(\frac{\mathbf{ID}}{\mathbf{L}}\right)^{0.33} \left(\frac{\mu}{\mu_{w}}\right)^{0.14}$$
(8-21)

For turbulent flow (Re_t > 10,000), h_i , can be expressed as:

$$\frac{\mathbf{h}_{i}.\mathrm{ID}}{\mathrm{k}} = \mathrm{C}\left(\frac{\mathrm{ID}.\mathrm{G}_{t}}{\mathrm{\mu}}\right)^{0.8} \left(\frac{\mathrm{c}\mathrm{\mu}}{\mathrm{k}}\right)^{0.33} \left(\frac{\mathrm{\mu}}{\mathrm{\mu}_{w}}\right)^{0.14}$$
(8-22)

where C = 0.021 for gases C = 0.023 for non-viscous liquids C = 0.027 for viscous liquids

For applications in turbulent flow, the viscosity ratio can be omitted, and Equation 8-22 reduces to:

$$h_i = C. k \left(\frac{12}{ID}\right) Re_t^{0.8} Pr_t^{0.33}$$
 (8-23)

for

 $0.7 < Pr_t < 17,000$

Figure 8-8 shows the heat transfer coefficient of fluids inside tubes. The heat transfer coefficient based on the outside of the tube, h_{io} , is:

$$\mathbf{h}_{io} = \mathbf{h}_{i} \left(\frac{\mathrm{ID}}{\mathrm{OD}} \right) \tag{8-24}$$

(9) Recently, Hedrick [12] developed a new correlation for the heat transfer coefficient in the transition region between laminar and turbulent flow. The equations for determining the inside film coefficient and based on the outside tube diameter, h_{io} , are:

$$\mathbf{h}_{io} = \left(\frac{16.1}{\text{OD}}\right) \left[\mathbf{B}_{i} \mathbf{k} \left(\frac{\mathbf{c}\boldsymbol{\mu}}{\mathbf{k}}\right)^{0.33} \left(\frac{\boldsymbol{\mu}}{\boldsymbol{\mu}_{w}}\right)^{0.14} \right]$$
(8-25)

where

$$B_{i} = (-3.08 + 3.075X + 0.32567X^{2} - 0.02185X^{3}) \times \left(\frac{10.\text{ID}}{\text{L}}\right)^{[1 - (X/10)^{0.256}]}$$

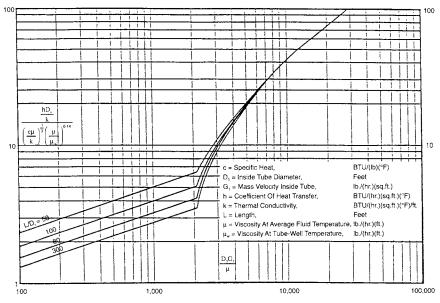


Figure 8-8. Coefficient of heat transfer fluids inside tubes.

and

$$X = \frac{Re_t}{1,000}$$

Simplified Equations

For common gases, Equation 8-23 can be simplified to give the approximate equation,

$$h_{i} = \frac{0.014C_{p}G_{i}^{0.8}}{(ID/12)^{0.2}}$$
(8-26)

Similarly for water at ordinary temperature and pressure,

$$\mathbf{h}_{i} = \frac{150(1+0.011t_{b})\mathbf{V}_{t}^{0.8}}{\mathrm{ID}^{0.2}}$$
(8-27)

where C_p = heat capacity of fluid, Btu/lb°F ID = tube inside diameter, inch G_t = mass velocity inside tube, lb/ft²hr t_b = average (i.e., bulk) temperature of water, °F $v_t^{0.8}$ = velocity of water, ft/sec

Total Pressure Drop, ΔPT

(10) From Re_t, determine the friction factor from Figure 8-9. Using the regression method in chapter one of the text, for Re_t \leq 1000, the friction factor correlation in Figure 8-9 can be approximated by

$$f_{t} = \frac{0.45}{Re_{t}^{0.9853}}$$
(8-28)

similarly, If $Re_1 > 1000$, the friction factor is approximated by

$$f_{t} = \frac{0.0028}{Re_{t}^{0.2514}}$$
(8-29)

(11) The fluid velocity, V_t , is:

$$V_{t} = \frac{G_{t}}{3600.\,\rho} \tag{8-30}$$

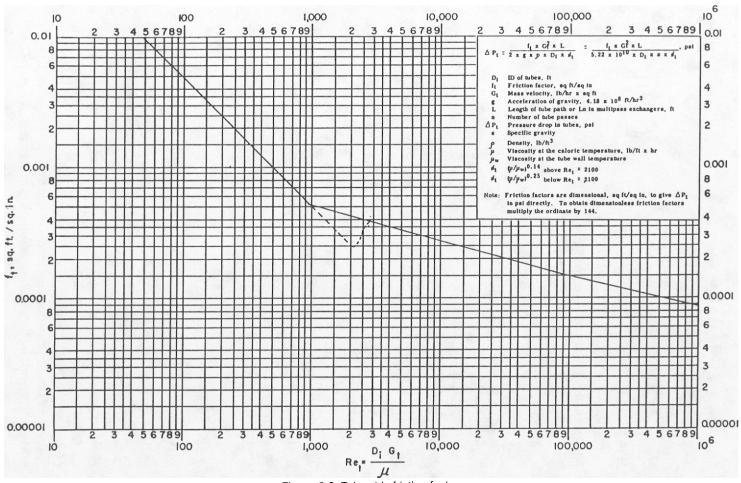


Figure 8-9. Tube side friction factors.

(12) Tube pressure drop, ΔP_{t} , psi

$$\Delta P_{t} = \frac{f_{t} \cdot G_{t}^{2} \cdot L \cdot n}{2g\rho(ID/12)}$$
(8-31)

where $G_t = lb/ft^2.s$

 $g = acceleration due to gravity, 32.17 ft/sec^2$

Alternatively, ΔP_t can be expressed as:

$$\Delta P_{t} = \frac{f_{t} \cdot G_{t}^{2} \cdot L \cdot n}{5.22 \times 10^{10} \cdot (ID/12) \cdot s \cdot \phi_{t}}$$
(8-32)

where $G_t = lb/ft^2hr$ and $\phi_t = \mu/\mu_w \approx 1$ s = specific gravity L = length of tube, ft n = number of tube passes

(13) The pressure drop of return losses, psi using Frank's [13] correlation is:

$$\Delta P_r = \frac{K.n.\rho.V_t^2}{2g(144)}$$
(8-33)

where K is the number of velocity heads loss per tube pass, where K = 2.5

(14) The pressure loss in both nozzles can be estimated from

$$\Delta P_{n} = 1.5 \rho \left(\frac{V_{n}^{2}}{2g}\right) \left(\frac{1}{144}\right)$$
(8-34)

(15) Total pressure drop ΔP_T , psi

$$\Delta P_{\rm T} = \Delta P_{\rm t} + \Delta P_{\rm r} + \Delta P_{\rm n} \tag{8-35}$$

If ΔP_t is unacceptable, it is necessary to assume a new pass arrangement and repeat the calculations. The heat transfer coefficient of the shell and tube heat exchanger is only acceptable if the design is within limits of the allowable pressure drop for the streams being heated or cooled. The selection of the optimum allowable pressure drop for an exchanger is a function of the overall process design.

Pressure drop calculations for more complex geometries of shell and tube exchangers, and for plate and spiral plate exchangers, are given by Sandler and Luckiewicz [14]. The acceptable range of pressure drops in a heat exchanger depends on the system operating pressure. Table 8-2 shows typical pressure drop ranges.

For high pressure drop design, fluid velocity may be the more limiting criterion, but a liquid velocity exceeding 15 ft/s can result in erosion-corrosion in an exchanger.

Shell Side

(1) Equivalent diameter, D_e , is calculated as follows:

Square pitch: $D_e = \frac{4 \times \text{free area}}{\text{wetted perimeter}}$

$$D_e = \frac{4(P_T^2 - \pi. OD^2/4)}{\pi. OD}$$
, inch (8-36)

Triangular pitch:

$$D_{e} = \left(\frac{1.72P_{T}^{2} - 0.5.\pi.OD^{2}}{0.5.\pi.OD}\right)$$
(8-37)

Clearance, C, between tubes is:

$$C = P_{T} - OD \tag{8-38}$$

Tube Arrangement

Tubes can be arranged in triangular, square, or rotated square pitch. Triangular tube layouts give better shell side heat transfer coefficients and provide more surface area for a given shell diameter. Alternatively, square pitch or rotated square pitch layouts are used when mechanical

Table 8-2
Acceptable Pressure Drop Range for
Shell and Tube Heat Exchanger

SYSTEM PRESSURE	PRESSURE DROP RANGE
Vacuum	5–10% of absolute system pressure.
1–10 psig	0.5–5 psi.
>10 psig	5 psi – 50% of system pressure.

cleaning of the outside of the tube is required. Widely spaced triangular patterns also allow for better cleaning. Both types of square pitches result in less shell-side turbulence and give low pressure drops, but lower heat transfer coefficients than triangular pitch.

Tables 8-3 to 8-10 show estimated tube counts for different square or triangular pitch arrangements. Figure 8-10 shows the tube layouts.

(2) Flow area a_s (assuming a plausible baffle spacing for the pressure drop allowed), is:

		TE Type				TEI Typ				ТЕІ Тур				TÉI Typ		
		Fix Tube:	ed sheet)utside Fioatin	Packed g Head			ins Floatin	ide g Head			U-Ti	ube	
Shell		No. of	Passes			No. of	Passes			No. of	Passes			No. of	Passes	
ID in.	1	2	4	6	1	2	4	6	1	2	4	6	2	4	6	8
5.047	22	20	16		19	18	12		19	14	12					
6.065	37	30	28		31	26	24		31	26	16					
7.981	68	66	60		61	52	48		56	52	44					
10.02	110	106	96		104	98	84		96	90	76					
12.00	170	164	148		151	142	128		151	138	128					
13.25	212	196	188		178	168	156		187	184	160					
15.25	283	270	252		241	232	220		258	242	224					
17.25	364	348	332		316	298	292		336	326	304					
19.25	454	440	420		396	388	352		421	412	392					
21.25	562	554	524		490	484	456		526	502	480					
23.25	668	646	612		588	570	548		608	598	556					
25.00																
27.00	922	902	868		812	808	764		868	836	804					
29.00																
31.00	1230	1212	1172		1106	1092	1040		1152	1124	1088					
33.00																
35.00	1590	1560	1516		1438	1430	1336		1496	1468	1424					

Table 8-3Tube Count for $^{3}/_{8}$ -in. OD Tubes on $^{13}/_{16}$ -in. Δ Pitch

Tube Count for 5/8 in. OD Tubes on 7/8 in. \triangle Pitch

		TEMA Type L or M				ТЕ Тур				TE Typ					MA e U	
			ed sheet			Outside Floatin	Packed g Head			ins Floatin	ide g Head			U-Т	ube	
Shell		No. of	Passes			No. of	Passes			No. of	Passes			No. of	Passes	
ID in.	1	2	4	6	1	2	4	6	1	2	4	6	2	4	6	B
5.047	22	18	16		19	14	12		14	14	12					
6.065	31	30	24		26	26	16		22	20	16					
7.981	61	52	48		55	48	44		51	48	40					
10.02	96	94	80		88	82	76		85	76	72					
12.00	151	138	132		130	124	112		130	120	112					
13.25	187	176	168		151	148	132		163	152	144					
15.25	241	232	224		206	196	184		216	214	196					
17.25	313	302	292		270	266	252		288	282	264					
19.25	396	384	352		336	334	312		358	350	340					
21.25	482	472	456		418	416	396		450	436	416					
23.25	568	554	536		506	492	472		526	506	484					
25.00																
27.00	792	780	752		704	700	660		724	720	696					
29.00																
31.00	1062	1030	1008		946	930	896		994	978	948					
33.00																
35.00	1356	1346	1304		1234	1220	1180		1288	1252	1220					

Courtesy of Tubular Exchanger Manufacturers Association (TEMA).

$$a_s = \frac{\text{ID. C. B}}{144. P_T}, \quad \text{ft}^2$$
 (8-39)

(3) Mass velocity, G_s is:

$$G_s = \frac{W}{a_s}, \quad \left(\frac{lb}{ft^2.hr}\right)$$
 (8-40)

Table 8-4Tube Count for $\frac{5}{8}$ -in. OD Tubes on $\frac{7}{8}$ -in.
 \Box Pitch

		TE Type I				TEI Typ				TÉ Typ				MA Ie U	
		Fix Tube:					Packed g Head		_	Ins Floatin			U-T	ube	
Shell		No. of	Passes			No. of	Passes			No. of	Passes		No. of	Passes	
ID in.	1	2	4	6	1	2	4	6	1	2	4	6	2	4	6
5.047	21	16	16		13	12	12		12	12	12				
5.065	26	26	24		22	22	16		21	16	16				
.981	52	52	52		45	44	44		45	38	32				
0.02	89	82	80		76	76	68		74	70	68				
2.00	128	124	120		109	104	92		109	92	88				
3.25	158	158	148		137	128	120		138	136	128				
5.25	213	208	208		177	176	164		188	184	176				
7.25	277	266	264		241	236	220		246	244	240				
9.25	344	332	336		293	284	284		316	308	304				
1.25	420	404	400		366	364	360		394	388	384				
23.25	502	476	488		436	432	432		456	448	444				
25.00															
27.00	694	668	664		612	608	608		640	636	624				
29.00															
31.00	922	910	908		828	812	812		862	848	840				
33.00															
35.00	1181	1166	1160		1070	1064	1048		1122	1112	1100				

		TEI Type L				ТЕ	MA He P			TE Typ				TEN Typ		
		Fix Tube:					Packed g Head			ins Floatin	ide g Head	_		U-TI	be	
Shell		No. of	Passes			No. of	Passes			No. of	Passes			No. of I	Passes	
ID in.	1	2	4	6	1	2	4	6	1	2	4	6	2	4	6	8
5.047	19	14	12		14	14	8		10	10	4		6	4		
6.065	27	26	20	18	22	18	16		19	18	12		14	8		
7.981	55	48	40	38	42	40	36	32	42	40	32	30	32	24		
10.02	85	76	72	68	74	72	60	54	73	66	60	54	56	52		
12.00	126	118	104	98	109	106	96	86	109	106	92	86	92	80		
13.25	151	148	140	134	130	124	112	108	140	138	124	108	114	104		
15.25	206	196	180	176	174	168	156	152	187	184	168	152	83	74		
17.25	268	266	240	230	241	222	216	210	253	242	224	210	220	204		
19.25	340	330	320	302	288	282	264	260	320	294	280	260	290	268		
21.25	416	408	392	384	384	368	344	338	400	380	352	338	360	340		
23.25	499	480	460	456	469	449	430	418	454	436	416	410	220	210		
25.00	576	558	530	516	544	529	500	490	514	498	471	465	506	488		
27.00	675	661	632	596	643	616	600	575	607	587	560	558	614	580		
29.00	790	773	736	720	744	732	704	695	707	690	663	657	720	684		
31.00	896	875	858		859	835	812	800	816	797	769	760	830	804		
33.00	1018	1011	976		973	959	926	900	931	910	876	870	944	916		
35.00	1166	1137	1098		1118	1093	1054	1010	1062	1039	998	993	1076	940		
37.00	1307	1277	1242		1253	1224	1184	1150	1200	1177	1135	1124	1218	1184		
39.00	1464	1425	1386		1392	1359	1318	1286	1341	1318	1282	1264	1366	1324		
42.00	1688	1669	1618		1616	1602	1552	1482	1558	1554	1502	1482	1600	1552		
45.00	1943	1912	1878		1870	1833	1800	1770	1875	1834	1736	1708	1854	1800		
48.00	2 2 29	2189	2134		2145	2107	2060	2025	2132	2100	1998	1964	2122	2064		
51.00	2513	2489	2432		2411	2395	2344	2305	2431	2392	2286	2250	2410	2356		
54.00	2823	2792	2752		2737	2683	2642	2612	2730	2684	2574	2536	2732	2668		
60.00	3527	3477	3414		3400	3359	3294	3220	3395	3346	3228	3196	3398	3336		

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(4) Shell velocity, V_s , ft/s is:

$$V_{s} = \frac{W}{3600. a_{s} \cdot \rho_{s}}$$
(8-41)

(5) Reynolds number, Re_s, is:

$$\operatorname{Re}_{s} = \frac{\operatorname{D}_{e} \cdot \operatorname{G}_{s}}{\mu}$$
(8-42)

Table 8-5Tube Count for $^{3}/_{4}$ -in. OD Tubes on 1-in. Δ Pitch

		TE Type	MA Lor M			TE Typ	MA De P			ТЕ Тур				ТЕМ Тур		
		Fix Tube	ied sheet			Dutside Floatin	Packed g Head	1		Ins Floatin				U-Tu	ube	
Shell		No. of	Passes			No. of	Passes			No. of	Passes	_		No. of F	asses	
ID in.	1	2	4	6	1	2	4	6	1	2	4	6	2	4	6	8
.047	14	14	12		10	10	8		10	10	4		6	4		
.065	22	20	16	22	19	18	16		19	14	12		10	8		
.981	42	40	36	40	38	36	32	28	37	32	28	26	28	24		
0.02	73	72	64	64	64	62	60	58	61	60	48	46	56	44		
2.00	109	86	80	98	96	94	84	78	96	94	80	78	86	72		
3.25	139	134	124	122	121	110	100	98	121	118	104	98	106	96		
5.25	187	180	168	164	151	146	140	138	163	164	144	140	148	136		
7.25	241	232	220	212	208	196	188	160	216	214	196	158	200	184		
9.25	296	290	280	270	258	242	232	230	276	270	260	235	254	240		
1.25	372	354	344	330	320	316	296	298	338	338	324	300	314	300		
3.25	434	420	404	404	380	372	364	335	396	396	376	339	388	368		
5.00	507	489	476	482	475	466	452	430	460	440	420	414	452	432		
7.00	604	594	568	582	530	526	508	495	558	554	536	494	538	524		
9.00	689	679	660	672	653	642	620	610	624	605	589	581	632	612		
1.00	808	804	772		724	696	688	669	756	744	716	669	732	708		
3.00	906	891	860		859	848	818	805	818	797	783	771	838	808		
5.00	1030	1026	1000		946	922	904	880	980	978	944	880	950	916		
7.00	1152	1134	1090		1106	1081	1054	996	1047	1039	1001	996	1074	1040		
9.00	1273	1259	1222		1218	1208	1174	1125	1172	1164	1130	1125	1200	1164		
2.00	1485	1461	1434		1426	1399	1376	1306	1367	1350	1322	1306	1406	1364		
5.00	1721	1693	1650		1652	1620	1586		1635	1608	1536	1504	1632	1584		
8.00	1968	1941	1902		1894	1861	1820		1887	1842	1768	1740	1870	1832		
1.00	2221	2187	2134		2142	2101	2060		2143	2104	2019	1992	2122	2076		
4.00	2502	2465	2414		2417	2379	2326		2399	2366	2270	2244	2396	2340		
0.00	3099	3069	3010		2990	2957	2906		2981	2940	2932	2800	2992	2936		

Tube Count for 3/4-in. OD Tubes on 1-in. 🗆 Pitch

	TEMA Type L or M					TEI Typ				TE! Typ			TEMA Type U				
		Fix Tubes				outside Floating		1		insi Floating			-	U-TL	ibe		
Shell		No. of I	Passes			No. of I	Passes			No. of I	Passes			No. of I	asses		
ID in.	1	2	4	6	1	2	4	6	1	2	4	6	2	4	6	8	
5.047	12	12	12		12	12	4		9	8	4	-	6	4			
6.065	21	16	16		16	16	12		16	16	12		8	8			
7.981	38	38	32		37	32	32	12	32	32	32	12	24	20			
10.02	61	60	52		57	56	52	24	56	52	52	24	44	40			
12.00	97	90	88		89	82	76	56	89	82	80	56	68	68			
13.25	117	116	112		97	94	88	80	104	104	96	80	90	88			
15.25	158	158	148		137	128	120	114	145	140	140	114	128	120			
17.25	216	208	188		177	176	164	160	188	184	180	160	176	168			
19.25	262	256	244		224	216	208	198	238	236	232	198	112	108			
21.25	316	316	308		274	270	268	260	304	292	284	260	138	134			
23.25	370	372	368		333	332	316	308	344	332	332	308	340	332			
25.00	442	432	428		414	406	392	344	398	386	366	344	400	388			
27.00	524	524	500		464	456	448	424	484	472	468	424	472	460			
29.00	602	596	580		570	562	548	496	554	532	510	496	554	544			
31.00	698	692	688		628	620	612	576	650	648	640	576	640	624			

(7) Prandtl number, Pr_s , is:

$$\Pr_{s} = \frac{c\mu}{k}$$
(8-43)

(8) The heat transfer coefficient, h_o , can be expressed as:

$$\frac{h_{o}.D_{e}}{k} = 0.36 \left(\frac{D_{e}G_{s}}{\mu}\right)^{0.55} \left(\frac{c\mu}{k}\right)^{0.33} \left(\frac{\mu}{\mu_{w}}\right)^{0.14}$$
(8-44)

Table 8-6
Tube Count for ³ / ₄ -in. OD Tubes on 1-in. Pitch

				Tu	be Cou	nt for	3/4-in. (OD Tul	oes on	1-in. (] Pitch	1					
			MA L or M			ТЕ: Тур				ТЕ Тур			TEMA Type U				
			red sheet				Packed g Head			Ins Floatin				U-Tu	be		
Sheil		No. of	Passes			No. of	Passes			No. of	Passes		No. of Passes				
ID in.	1	2	4	6	1	2	4	6	1	2	4	6	2	4	6	8	
33.00	782	768	768		742	732	732	668	730	712	682	668	724	720			
35.00	894	892	880		816	812	804	760	848	828	824	760	836	812			
37.00	1004	978	964		952	931	928	872	937	918	882	872	9 40	924			
39.00	1102	1096	1076		1062	1045	1026	972	1048	1028	996	972	1048	1040			
42.00	1283	1285	1270		1232	1222	1218	1140	1224	1200	1170	1140	1222	1204			
45.00	1484	1472	1456		1424	1415	1386	1336	1421	1394	1350	1336	1420	1400			
48.00	1701	1691	1670		1636	1634	1602	1536	1628	1598	1548	1536	1624	1604			
51.00	1928	1904	1888		1845	1832	1818	1764	1862	1823	1779	1764	1852	1820			
54.00	2154	2138	2106		2080	2066	2044	1992	2096	2048	2010	1992	2084	2064			
60.00	2683	2650	2636		2582	2566	2556	2476	2585	2552	2512	2476	2596	2564			

Tube Count for ¾-in. OD Tubes on 1-in.	Pitch
--	-------

		TE Type I	MA L or M			ТЕ Тур					MA ke S			TEI Typ		
		Fix Tube:	ed sheet	_			Packed g Head	1		ins Floatin	ide g Head			U-T(ibe	
Shell		No. of	Passes			No. of	Passes	_		No. of	Passes			No. of I	asses	
ID in.	1	2	4	6	1	2	4	6	1	2	4	6	2	4	6	8
5.047	12	10	8		12	10	8						4	4		
6.065	21	18	16		16	12	8						10	8		
7.981	37	32	28		32	28	24						24	20		
10.02	61	54	48		52	46	40						42	36		
12.00	97	90	84		81	74	68						66	64		
13.25	113	108	104		97	92	84						86	80		
15.25	156	146	136		140	134	128						124	116		
17.25	208	196	184		188	178	168						174	164		
19.25	256	244	236		241	228	216						218	202		
21.25	314	299	294		300	286	272						272	260		
23.25	379	363	352		359	343	328						334	320		
25.00	448	432	416		421	404	392						390	380		
27.00	522	504	486		489	472	456						468	452		
29.00	603	583	568		575	556	540						550	532		
31.00	688	667	654		660	639	624						626	608		
33.00	788	770	756		749	728	708						720	700		
35.00	897	873	850		849	826	804						818	796		
37.00	1009	983	958		952	928	908						928	904		
39.00	1118	1092	1066		1068	1041	1016						1036	1016		
42.00	1298	1269	1250		1238	1216	1196						1220	1192		
45.00	1500	1470	1440		1432	1407	1378						1412	1384		
48.00	1714	1681	1650		1644	1611	1580						804	788		
51.00	1939	1903	1868		1864	1837	1804						1834	1804		
54.00	2173	2135	2098		2098	2062	2026						2072	2036		
60.00	2692	2651	2612		2600	2560	2520		_				2584	2544		

For turbulent flow, the viscosity ratio can be omitted, and Equation 8-44 becomes:

$$h_o = 0.36 \frac{k}{D_c} Re_s^{0.55} Pr_s^{0.33}$$
 (8-45)

Figure 8-11 shows the shell side heat transfer coefficient as a function of the Reynolds number.

		TEI Type I				ТЕ! Тур				те: Тур				ТЕЛ		
		Fix Tubes				Dutside Floatin	Packed g Head	1		ins Floatin				U-T.	ibe	
Sheli		No. of	Passes			No. of	Passes			No. of	Passes			No. of I	asses	
ID in.	1	2	4	6	1	2	4	6	1	2	4	6	2	4	6	6
5.047	8	6	4		7	4	4						0	0		
5.065	14	14	8		10	10	4						4	4		
7.981	26	26	16		22	18	16	14	18	14	8	12	14	8		
10.02	42	40	36		38	36	28	24	33	28	16	18	26	24		
12.00	64	61	56		56	52	48	46	51	48	42	44	44	36		
3.25	85	76	72		73	72	60	44	73	68	52	44	56	52		
5.25	110	106	100		100	98	88	80	93	90	78	76	86	76		
7.25	147	138	128		130	126	116	104	126	122	112	102	114	104		
9.25	184	175	168		170	162	148	140	159	152	132	136	152	136		
21.25	227	220	212		212	201	188	176	202	192	182	172	192	176		
23.25	280	265	252		258	250	232	220	249	238	216	212	232	220		
25.00	316	313	294		296	294	276	250	291	278	250	240	270	256		
27.00	371	370	358		355	346	328	300	345	330	298	288	322	304		
29.00	434	424	408		416	408	392	360	400	388	356	348	378	364		
31.00	503	489	468		475	466	446	420	459	450	414	400	444	424		
33.00	576	558	534		544	529	510	498	526	514	484	464	508	492		
35.00	643	634	604		619	604	582	566	596	584	548	536	578	560		
37.00	738	709	684		696	679	660	646	672	668	626	608	660	632		
39.00	804	787	772		768	753	730	723	756	736	704	692	740	712		
2.00	946	928	898		908	891	860	840	890	878	834	808	872	836		
15.00	1087	1069	1042		1041	1017	990	968	1035	1008	966	948	1010	980		
18.00	1240	1230	1198		1189	1182	1152	1132	1181	1162	1118	1092	1156	1124		
51.00	1397	1389	1354		1348	1337	1300	1280	1350	1327	1277	1254	1322	1284		
4.00	1592	1561	1530		1531	1503	1462	1440	1520	1492	1436	1416	1496	1452		
50.00	1969	1945	1904		1906	1879	1842	1802	1884	1858	1800	1764	1866	1828		

Table 8-7
Tube Count for 1-in. OD Tubes on $1^{1/4}$ -in. Δ Pitch

Tube Count for 1-in. OD Tubes on 11/4-in. \Diamond Pitch

		TEN Type L				TEN Type					MA be S			ТЕМ Турн		
		Fix Tubes				utside Floating	Packed Head				ide Ig Head			U-TI	ıbe	
Shell		No. of I	Passes			No. of I	Passes			No. of	Passes			No. of I	asses	
ID in.	1	2	4	6	1	2	4	6	1	2	4	6	2	4	6	8
5.047	8	6	4		5	4	4						0	0		
6.065	12	10	8		12	10	8						4	4		
7.981	24	20	16		21	18	16						10	8		
10.02	37	32	28		32	32	28						24	10		
12.00	57	53	48		52	46	40						36	32		
13.25	70	70	64		61	58	56						50	44		
15.25	97	90	84		89	82	76						70	64		
17.25	129	120	112		113	112	104						96	88		
19.25	162	152	142		148	138	128						124	120		
21.25	205	193	184		180	174	168						156	152		
23.25	238	228	220		221	210	200						200	188		
25.00	275	264	256		261	248	236						232	220		
27.00	330	315	300		308	296	286						282	268		
29.00	379	363	360		359	345	336						330	320		
31.00	436	422	410		418	401	388						382	368		

Pressure Drop (ΔP)

(9) From the fluid Reynolds number, Re_{s} , obtain the friction factor, f_{s} , from Figure 8-12. A regression method of Figure 8-12 shows that for $\text{Re}_{s} < 500$,

$$f_s = \frac{0.11183}{Re_s^{0.59246}}$$
(8-46)

Table 8-8Tube Count for 1-in. OD Tubes on 11/4-in.

				Tul	be Cou	nt for	1-in. O[7 Tub	es on 1	1 ¼-in.	◊ Pitc	h				
		TE	MA			TE	MA			TE	MA			TER	đΑ.	
		Туре	Lor Mi			Тур	e P			Ту	be S			Тур	eυ	
		Fixed Tubesheet No. of Passes				Dutside Floatin	Packed g Head				side Ig Head			U-Ti	ibe -	
Shell						No. of	Passes			No. of	Passes			No. of I	asses	
lD in.	1	2	4	6	1	2	4	6	1	2	4	6	2	4	6	8
33.00	495	478	472		477	460	448						440	424		
35.00	556	552	538		540	526	508						498	484		
37.00	632	613	598		608	588	568						562	548		
39.00	705	685	672		674	654	640						630	620		
42.00	822	799	786		788	765	756						744	728		
45.00	946	922	912		910	885	866						872	852		
48.00	1079	1061	1052		1037	1018	1000						1002	980		
51.00	1220	1159	1176		1181	1160	1142						1138	1116		
54.00	1389	1359	1330		1337	1307	1292						1292	1264		
60.00	1714	1691	1664		1658	1626	1594						1604	1576		

Tube Count for 1-in. OD Tubes on 11/4-in. 🗆 Pitch

			MA Lor M				MA be P				MA De S			ТЕІ Тур		
			sheet				g Head				ide g Head			U-Ti	ube	
Shell		No. of	Passes			No. of	Passes			No. of	Passes			No. of	Passes	
ID in.	1	2	4	6	1	2	4	6	1	2	4	6	2	4	6	8
5.047	9	6	4		5	4	4	-	5	4	4	_	0	0		
6.065	12	12	12		12	6	4	-	12	6	4	-	4	4		
7.981	22	20	16		21	16	16	12	17	12	8	12	12	8		
10.02	38	38	32		32	32	32	18	30	30	16	18	24	20		
12.00	56	56	52		52	52	44	24	52	48	42	24	38	36		
13.25	69	66	66		61	60	52	50	61	56	52	50	52	48		
15.25	97	90	88		89	84	80	64	85	78	62	64	72	68		
17.25	129	124	120		113	112	112	96	108	108	104	96	98	96		
19.25	164	158	148		148	144	140	114	144	136	130	114	128	124		
21.25	202	191	184		178	178	172	156	173	166	154	156	166	156		
23.25	234	234	222		216	216	208	192	217	208	194	192	200	196		
25.00	272	267	264		258	256	256	212	252	240	230	212	240	232		
27.00	328	317	310		302	300	296	260	296	280	270	260	284	276		
29.00	378	370	370		356	353	338	314	345	336	310	314	332	332		
31.00	434	428	428		414	406	392	368	402	390	366	368	290	384		
33.00	496	484	484		476	460	460	420	461	452	432	420	442	436		
35.00	554	553	532		542	530	518	484	520	514	494	484	254	248		
37.00	628	621	608		602	596	580	550	588	572	562	548	574	560		
39.00	708	682	682		676	649	648	625	661	640	624	620	644	628		
42.00	811	811	804		782	780	768	730	776	756	738	724	758	748		
45.00	940	931	918		904	894	874	850	900	882	862	844	872	868		
48.00	1076	1061	1040		1034	1027	1012	980	1029	1016	984	972	1002	988		
51.00	1218	1202	1192		1178	1155	1150	1125	1170	1156	1126	1114	1146	1140		
54.00	1370	1354	1350		1322	1307	1284	1262	1310	1296	1268	1256	1300	1288		
60.00	1701	1699	1684		1654	1640	1632	1585	1641	1624	1598	1576	1620	1604		

for $\text{Re}_{s} > 500$,

$$f_s = \frac{0.1159}{Re_s^{0.18597}} \tag{8-47}$$

The shell side pressure drop, ΔP_s , is:

$$\Delta P_{s} = \left\{ \frac{f_{s} \cdot G_{s}^{2} \cdot (ID/12) \cdot (N+1)}{2 \cdot g \cdot \rho \cdot D_{e} \phi_{s}} \right\} (N_{s})$$
(8-48)

Table 8-9 Tube Count for $1^{1}/_{4}$ -in. OD Tubes on $1^{9}/_{16}$ -in. Δ Pitch

		TEI Type I				TEI Typ					MA De S			ТЕМ Туре		
		Fix Tubes				outside Floating	Packec Head				side 1g Head			U-Tu	ibe	
Shell		No. of	Passes			No. of I	Passes			No. of	Passes			No. of F	basses	
ID in.	1	2	4	6	1	2	4	6	1	2	4	6	2	4	6	8
5.047	7	4	4		0	0	0	-					0	0		
6.065	8	6	4		7	6	4	-					0	0		
7.981	19	14	12		14	14	8	-					6	4		
10.02	29	26	20		22	20	16	-					14	12		
12.00	42	38	34		37	36	28	22					22	20		
13.25	52	48	44		44	44	36	28					32	28		
15.25	69	68	60		64	62	48	45					48	44		
17.25	92	84	78		85	78	72	69					64	60		
19.25	121	110	104		09	102	96	86					86	80		
21.25	147	138	128		30	130	116	112					114	104		
23.25	174	165	156		63	152	144	130					138	132		
25.00	196	196	184		84	184	172	164					162	152		
27.00	237	226	224		21	216	208	196					196	184		
29.00	280	269	256		62	252	242	228					232	220		
31.00	313	313	294		02	302	280	270					268	256		
33.00	357	346	332		45	332	318	305					310	296		
35.00	416	401	386		92	383	364	357					356	344		
37.00	461	453	432		42	429	412	407					404	388		
39.00	511	493	478		93	479	460	449					452	440		
42.00	596	579	570		76	557	544	512					534	522		
45.00	687	673	662		57	640	628	596					626	612		
48.00	790	782	758		56	745	728	696					720	700		
51.00	896	871	860		59	839	832	820					822	800		
54.00	1008	994	968		64	959	940	892					930	908		
60.00	1243	1243	1210	11	99	1195	1170	1116					1160	1140		

Tube Count for 11/4-in. OD Tubes on 11/16-in. 🗆 Pitch

		TEN Type L				TEN Type				ТЕМ Тур				TEP Typ		
	Fixed Tubesheet No. of Passes					utside loating	Packed Head			Insi Floating				U-TI	ipe	
Shell		No. of F	asses		1	No. of F	asses			No. of F	Passes			No. of I	Passes	
ID in.	1	2	4	6	1	2	4	6	1	2	4	6	2	4	6	8
5.047	4	4	4		0	0	0	0	0	0	0	0	0	0		
6.065	6	6	4		6	6	4	0	6	6	4	0	0	0		
7.981	12	12	12		12	12	12	0	12	12	4	0	6	4		
10.02	24	$\begin{array}{cccccccccccccccccccccccccccccccccccc$			21	16	16	12	21	12	8	12	12	8		
12.00	37	34	32		32	32	32	18	29	28	16	18	20	20		
13.25	45	42	42		38	38	32	24	38	34	34	24	28	28		
15.25	61	60	52		52	52	52	48	52	48	44	48	42	36		
17.25	80	76	76		70	70	68	50	70	66	56	50	56	56		
19.25	97	95	88		89	88	88	80	85	84	70	80	74	68		
21.25	124	124	120		112	112	112	96	108	108	100	96	98	96		
23.25	145	145	144		138	138	130	114	136	128	128	114	124	120		
25.00	172	168	164		164	164	156	136	154	154	142	136	140	136		

where $G_s = lb/ft^2hr$ and $f_s = \mu/\mu_w \approx 1$ $g = acceleration due to gravity, <math>4.18 \times 10^8$ ft/hr² $\rho = density, lb/ft^3$ $N_s = number of shells per unit$

Alternatively, ΔP_s can be expressed as:

$$\Delta P_{s} = \left\{ \frac{f_{s} \cdot G_{s}^{2} \cdot (ID/12) \cdot (N+1)}{5.22 \times 10^{10} \cdot D_{e} \cdot s \cdot \phi_{s}} \right\} (N_{s})$$
(8-49)

Table 8-10Tube Count for $1^{1/4}$ -in. OD Tubes on $1^{9/16}$ -in. \Box Pitch

		TEN Type L				те! Тур				тем Тур				те! Тур		
		Fix Tubes				outside Floating	Packed Head			Insi Floating				U-Ti	ibe	
Sheli		No. of I	Passes			No. of I	Passes			No. of I	Passes			No. of I	Passes	
ID in.	1	2	4	6	1	2	4	6	1	2	4	6	2	4	6	8
27.00	210	202	202		193	184	184	172	184	180	158	172	176	176		
29.00	241	234	230		224	224	216	198	217	212	204	198	200	196		
31.00	272	268	268		258	256	256	236	252	248	234	236	232	232		
33.00	310	306	302		296	296	282	264	289	276	270	264	272	268		
35.00	356	353	338		336	332	332	304	329	316	310	304	312	296		
37.00	396	387	384		378	370	370	358	372	368	354	340	348	348		
39.00	442	438	434		428	426	414	408	420	402	402	392	396	392		
42.00	518	518	502		492	492	484	464	485	476	468	464	472	456		
45.00	602	602	588		570	566	556	544	565	554	546	544	552	536		
48.00	682	681	676		658	648	648	620	653	636	628	620	628	620		
51.00	770	760	756		742	729	722	712	738	726	720	705	712	708		
54.00	862	860	856		838	823	810	804	837	820	812	804	808	804		
60.00	1084	1070	1054		1042	1034	1026	1008	1036	1028	1012	1008	1012	992		

Tube Count for 11/4-in. OD Tubes on 19/16-in. ◇ Pitch

		TEN Type L				ТЕМ Тур		-		ТЕ Тур	MA Ie S			ТЕЛ Тур		
		Fix Tubes				utside Floating	Packed Head			ins Fioatin				U-Ta	ube	
Shell		No. of	Passes		1	No. of I	Passes			No. of	Passes			No. of i	Passes	
ID in.	1	2	4	6	1	2	4	6	1	2	4	6	2	4	6	8
5.047	5	4	4		0	0	0						0	0		
6.065	6	6	4		5	4	4						0	0		
7.981	13	10	8		12	10	8						4	4		
10.02	24	20	16		21	18	16						12	12		
12.00	37	32	28		32	28	28						20	20		
13.25	45	40	40		37	34	32						26	24		
15.25	60	56	56		52	52	48						40	36		
17.25	79	76	76		70	70	64						56	52		
19.25	97	94	94		90	90	84						74	68		
21.25	124	116	112		112	108	104						96	88		
23.25	148	142	136		140	138	128						120	112		
25.00	174	166	160		162	162	156						142	136		
27.00	209	202	192		191	188	184						170	164		
29.00	238	232	232		442	430	416						200	192		
31.00	275	264	264		261	249	244						228	220		
33.00	314	307	300		300	286	280						268	256		
35.00	359	345	334		341	330	320						306	296		
37.00	401	387	380		384	372	360						346	336		
39.00	442	427	424		428	412	404						390	380		
42.00	522	506	500		497	484	472						456	448		
45.00	603	583	572		575	562	552						542	528		
48.00	682	669	660		660	648	640						618	604		
51.00	777	762	756		743	728	716						708	692		
54.00	875	857	850		843	822	812						802	784		
60.00	1088	1080	1058		1049	1029	1016						1010	984		

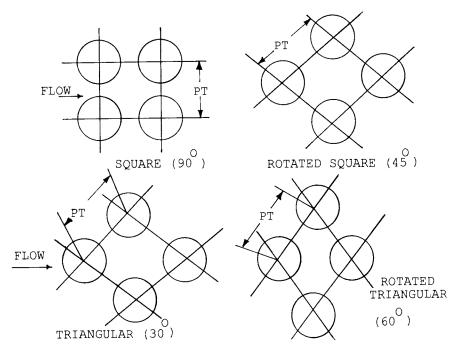


Figure 8-10. Tube layouts.

where
$$G_s = lb/ft^2hr$$
 and $\phi_t = \mu/\mu_w \approx 1$
s = specific gravity
N = number of baffles
N_s = number of shells per unit.

If ΔP_s is unacceptable, assume a new baffle spacing.

(10) The overall heat transfer coefficient, U, is:

$$U = \frac{1}{\frac{1}{h_{io}} + \frac{1}{h_o} + F_T + F_s}, \quad \frac{Btu}{hr ft^2 \circ F}$$
(8-50)

(11) The outside area of the heat exchanger unit A, ft^2 can be expressed as:

$$\mathbf{A} = \left(\frac{\mathbf{OD}_{t}}{12} \cdot \boldsymbol{\pi} \cdot \mathbf{n} \cdot \mathbf{N}_{s} \cdot \mathbf{L}\right)$$
(8-51)

(12) Heat load on unit Q, Btu/hr, is:

$$Q = U.A.\Delta t \tag{8-52}$$

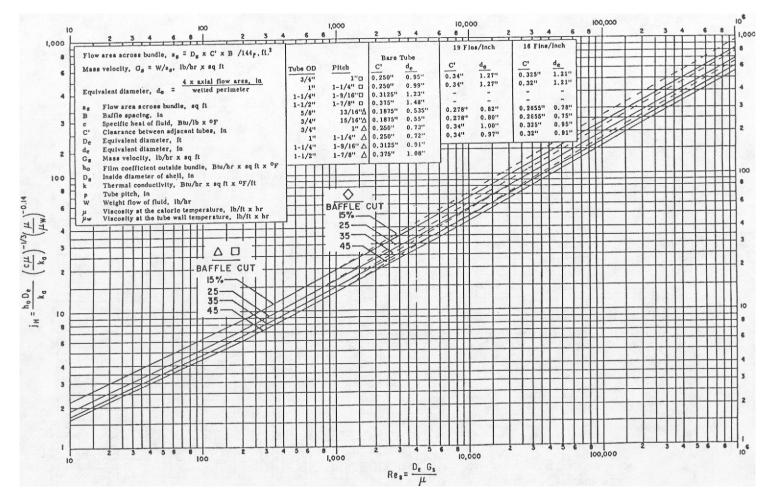


Figure 8-11. Shell-side heat transfer parameter J_H versus N_{Re} for various baffle cuts.

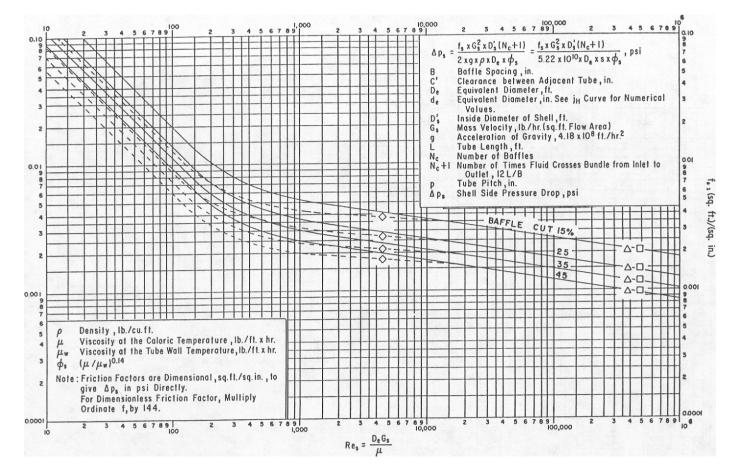


Figure 8-12. Friction factors for shell-side surfaces of low-finned and bare tubes.

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Fouling

The scale or fouling resistances represent a necessary safety factor that increases the surface of the heat exchanger. This enables the full process duty requirements to be attained between cleaning periods. When an exchanger is first placed in operation there is no dirt or scale on the tubes; consequently, the overall resistance consists of the two film and the tube wall resistances. During operation, dirt or scale accumulates on the surface of the tubes and the overall heat transfer rate decreases as the dirt buildup increases. The rate of this scale or dirt depends on the cleanliness or fouling tendencies of the process fluids.

The fouling experienced in most CPI operations are due to:

Crystallization. When salts having an inverse solubility characteristic precipitate on a heat transfer surface hotter than the flowing fluid.

Sedimentation. Caused by deposits of particulate matter such as clay, sand, or rust.

Corrosion. When the heat transfer surface material reacts with the fluid to yield corrosion deposits.

Chemical reaction product buildup. Caused by organic products and polymers. The surface temperature and presence of reactants, particularly oxygen, can have a very significant effect on the rate of accumulation.

Coking. On high temperature surface results in hydrocarbon deposits.

Biofouling. Deposits formed when biological mechanisms attach and grow on the heat transfer surface. Untreated cooling water systems are particularly susceptible to biofouling.

The range of overall heat transfer coefficients (U) is about 10 - 200 Btu/hr ft^{2°}F. Table 8-11 lists the U-values for various types of equipment, and fouling factors of some flowing media are shown in Table 8-12. Table 8-13 illustrates a heat exhanger specification form. Factors governing the selection of process fluids in the tube and shell sides of an exchanger are illustrated in Table 8-14.

Nomenclature

A = outside area of the heat exchanger unit, ft^2 a_t = flow area, ft^2

Equipment	Process	U
Shell-and-tube exchanger	gas (latm)–gas (latm)	1–6
	gas (250atm)–gas (250atm)	25-50
	liquid–gas (1atm)	2-12
	liquid–gas (250atm)	35-70
	liquid–liquid	25-200
	liquid-condensing vapor	50-200
Double-pipe exchanger	gas (latm)–gas (latm)	26
	gas (250atm)–gas (250atm)	25-90
	liquid–gas (250atm)	35-100
	liquid–liquid	50-250
Irrigated tube bank	water–gas (1atm)	3-10
	water–gas (250atm)	25-60
	water-liquid	50-160
	water-condensing vapor	50-200
Plate exchanger	water-gas (1atm)	3-10
	water-liquid	60–200
Spiral exchanger	liquid–liquid	120-440
liquid-condensing steam		160-600
Compact exchanger	gas (1atm)–gas (1atm)	2-6
	gas (1atm)–liquid	3–10
Stirred tank, jacketed	liquid-condensing steam	90–260
	boiling liquid-condensing steam	120-300
	water-liquid	25-60
Stirred tank, coil inside	liquid-condensing steam	120440
	water-liquid	90–210

Table 8-11Ranges of Overall Heat Transfrer Coefficientsin Various Types of Exchangers[U Btu/hrft2°F]

Source: S. M. Walas, Chemical Process Equipment—Selection and Design, Butterworths Series in Chemical Engineering.

Table 8-12Recommended Minimum Fouling Resistances

Gases and Vapors	Fouling Factor
Centrifugal Compressor exhaust	0.001
Reciprocating Compressor exhaust	0.01
Reciprocating Compressor refrigerant vapor	0.0025
Centrifugal compressor refrigerant vapor	0.0015
Oil-free and clean high-quality steam	0.0003
Oil-free and clean low-quality steam	0.0005
Oil-bearing steam	0.001
Compressed air	0.002
Acid gas	0.001
Solvent vapors	0.001
Natural gas	0.001
Liquids	
Bay water	0.0025
Distilled water	0.0005
Hard well water	0.0033
Untreated cooling tower water	0.0033
Treated cooling tower water	0.0015
Engine jacket water	0.0012
Treated boiler feed water	0.0015
Fuel oils	0.006
Clean organic solvents	0.001
Vegetable oils	0.004
Refrigerant liquids	0.001
Industrial heat transfer oils	0.001
Hydraulic fluid	0.001
Natural gasoline and liquefied petroleum gases	0.001
Rich oil	0.001
Lean oil	0.002

Table 8-13 Heat Exchanger Specification Sheet (Courtesy of Procede)

Hee	it Exc	hano	er d	late	l cl	ieet		ļ	Equipment No	(Tag)			
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SIZE			TYPE		0000				F UNITS			+	3
SHELLS PER UNIT	· ·			ICE PER		ECTED IN (parailel or ser	nes)				4+	4
SURFACE FER UNIT	L.,,						L.						6
		P	erforn	nanc	e of	one Uni	t						7
				SHELL	SIDE	·			TUBE S	DE			8
FLUID CIRCULATING		+											9
TOTAL FLUID ENTERING		<u> </u>	-								OUT		10
VAPOUR		+	IN	-+		OUT		IN			001		1
LIQUID		1 1										\mathbf{t}	13
STEAM					-							+	14
WATER													1
NON-CONDENSABLES		1											10
FLUID VAPOURISED OR CO							┶╋┸			<u> </u>		1	1
SPECIFIC GRAVITY LIQUID Mol W1 VAPOUR		+	• •	· ··									11
Mol WI VAPOUR Mol WI NON-CONDENSABL	ES	+											20
VISCOSITY LIQUID		1											21
LATENT HEAT		1											2
SPECIFIC HEAT								•					2
THERMAL CONDUCTIVITY		· · ·	1	· · · · · ·									24
TEMPERATURE				1.1			┶╋╍┖					1	2:
OPERATING PRESSURE		+											2
VELOCITY No. OF PASSES		+											2
PRESSURE DROP		ALLOW	1	Т	CALC.		ALL	ow	· · · · · ·	CALC.		T	29
FOULING RESISTANCE						[-			Ţ		-	30
HEAT EXCHANGED						MT	D (CORREC	TED)					31
TRANSFER RATE SERVICE						<u> </u>	EAN						32
		C	onstru	uctio	1 of (one Shel	1						31
DESIGN PRESSURE						1							3/
TEST PRESSURE							· · · · · · · · · · · · · · · · · · ·						3
DESIGN TEMPERATURE													3
METAL TEMPERATURE						l							3
TUBES			No. OE		T	ICKNESS	LE	NGTH		т			3
SHELL			1.D.						L COVER				4
CHANNEL TUBE SHEET STATIONARY			CHAN	NEL CO	VER	l		FLIN	IG HEAD COV	VER		+-1	4
BAFFLES CROSS			TYPE						ING %CUI			1-1	4
TUBE SUPPORTS			TYPE						ING				4
LONG BAFFLE			TYPE					SEAL	4				4
IMPINGEMENT BAFFLE			TYPE						. STRIPS			+	4
TYPE OF JOINT			TUBE				+		E ATTACHM	r		+	4
GASKETS SHELL IN			CHAN						ATING HEAD			+-+	4
CONNECTIONS SHELL IN	N -		INTER						LOUT			+ - + - + - + - + - + - + - + - + - + -	4
CONNECTIONS CHANNEL I CORROSION ALLOWABLE :			AUTICK	CONIN					NNEL OUT			+	5
EXPANSION BELLOWS			BOLTS	;				NUT				+ 1	5
DESIGN CODE			X-RAY					S.R.					5
INSPECTION			PAINT	ING	-				LATION				5
WEIGHT OF ONE UNIT EM	ny 🗌		OPERA						E OF ENQUIR	Y		11	5
DATE OF ORDER			ORDEI	R No.			l_	DRG	NO.			+	5
INSPECTION FITTING ATT	CHMENT BY	T	3	1		MANUFAC	URER	6	· · · · · · · · ·		· ···	4	5
Prepared Checked			2	+	-+-			6 5	+				5
Approved			$+\frac{2}{1}$	1	-			4	+ +			-	6
	Engineering	Process	REV	By		Appr.	Date	REV	By	Appr.	Dat	ic	6
Service				pany				Addr		- FRA			6
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Equipment No.			-										6
Project No.			1										16

Table 8-14Factors Which Influence the Selection ProcessFluids in a Shell and Tube Heat Exchanger

Tube Side	Shell Side
CORROSION: When special alloy materials are necessary to minimize corrosion. It is more economical to place the corrosive fluids in the tubes. Only the tubes, tube sheets and channel need to be alloy. In comparison, if the shell side was selected, the shell side and baffles would have to be alloy in addition to the tubes and tube sheets.	CONDENSING SERVICE: The large free area on the shell side permits minimum pressure drop arrangements as well as higher condensate loadings.
EXCESSIVE FOULING: If mechanical cleaning is desired, the fouling medium should be placed in the tube side. This enables the tubes to be cleaned without removing the bundle from the shell.	BOILING SERVICE: Vapor disen- gaging space may be provided on the shell side which eliminates the need for a separate dry drum.
HIGH TEMPERATURE: If high temperature service requires special alloy construction, the cost of an alloy shell and bonnet will be saved by using the tube side.	CRITICAL PRESSURE DROP: By varying the baffle, spacing, very low pressure drops may be attained.
HIGH PRESSURE: Tubes, tube sheets, and channel need be designed only for high pressure if the tube side is utilized. Also, integral tube sheet and channel design may be used, thus eliminating one gasketed joint.	VISCOUS FLUIDS: If mechanical cleaning is not required, higher heat transfer rates may be obtained by placing the viscous fluid on the shell side. Due to the flow pattern across the tube bank, turbulent flow may be maintained on the shell side at mass velocities which would yield laminar flow on the tube side.

Source: Morton [7]

(text continued from page 623)

 $a_s = flow area, ft^2$ B = baffle spacing, inchC = clearance between adjacent tube, inch D_e = equivalent diameter, ft F_c = temperature difference correction factor, $\Delta t = (F_c)(LMTD)$ F_T = tube side fouling factor, hr ft²°F/Btu F_s = shell side fouling factor, hr ft²°F/Btu f_{i} = tube side friction factor (ft²/in²) $f_s =$ shell side friction factor (ft²/in²) G_t = tube side mass velocity, (lb/ft²hr) or (lb/ft²sec) G_s = shell side mass velocity, (lb/ft²hr) g = acceleration due to gravity, 32.17 ft/sec² (4.18×10^8 ft/hr²) h_i , h_o = heat transfer coefficient for inside and outside fluids, respectively, Btu/hrft2°F h_{io} = value of hi when referred to the tube outside diameter, ID = inside diameter, inch (ft) $j_{\rm H}$ = factor for heat transfer, dimensionless. k = thermal conductivity, Btu/hrft°FL = tube length, ftLMTD = log mean temperature difference, °F N = number of shell side baffles $N_s =$ number of shells per unit $N_i =$ number of tubes n = number of tube passesOD = outside diameter, inchP = temperature group $(t_2 - t_1)/(T_1 - t_1)$ P_r = prandtl number, dimensionless P_{t} = tube pitch, inch $\Delta P_{T}, \Delta P_{t} = \text{total}, \text{ tube side pressure drop, psi.}$ ΔP_r , ΔP_n = return and nozzle side pressure drops, psi ΔP_s = shell side pressure drop, psi Q = heat flow, Btu/hr

R = temperature group $(T_1 - T_2)/(t_2 - t_1)$, dimensionless

Re = reynolds number, dimensionless.

 R_d = combined dirt factor, hr ft²°F/Btu

 T_1 = hot fluid inlet temperature, °F

 T_2 = hot fluid outlet temperature, °F

 $t_1 = cold fluid inlet temperature, °F$

 $t_2 = cold fluid outlet temperature, °F$

 Δt = true temperature difference, °F

U = overall heat transfer coefficient, Btu/hrft^{2°}F V_n = nozzle side fluid velocity, ft/s V_s = shell side fluid velocity, ft/s V_t = tube side fluid velocity, ft/s W = mass flow rate of hot fluid, lb/hr w = mass flow rate of cold fluid, lb/hr ϕ = viscosity ratio, (μ/μ_w)^{0.14} μ = fluid viscosity, cP × 2.42 = lb/ft. hr. ρ = fluid density, lb/ft³

Subscripts

s = shellt = tube

DOUBLE PIPE HEAT EXCHANGER

A double pipe exchanger consists of one or more pipes or tubes inside a pipe shell. Basically two straight pipe lengths are connected at one end to form a U or "hair-pin." Longitudinal fins may be used on the outside of the inner tube. The flow is true countercurrent, which can be beneficial when very close temperature approaches or very long temperature ranges are required. Such an exchanger is used in service when the heat duty is moderate (that is, UA < 100,00 Btu/hr°F), or when one stream is a viscous liquid, or when flow rates are small. This type of exchanger is suitable for high pressure applications because of their smaller diameters. Also, several hairpin sections provide for flexibility in matching heat exchanger requirements with changing process conditions. A double pipe exchanger is suited for "dirty" service because it is easy to dismantle and clean. In addition, a pipe exchanger can be considered in the following situations:

- 1. When the shell side coefficient is less than half that of the tube side; the annular side coefficient can be made comparable to the tube side.
- 2. When a high pressure can be catered for more economically in the annulus than in a larger diameter shell.
- 3. At duties requiring 100 200 ft² of the surface, which make it more economical.
- 4. When a true countercurrent flow can be obtained, thus eliminating temperature crosses that require multishell shell and tube units.

The equations required for the rating of a double pipe exchanger are as follows.

The Equations

Process Conditions Required

Hot fluid: T_1 , T_2 , W, C, μ , k, ΔP , R_D , s or ρ

Cold fluid: $t_1, t_2, w, c, \mu, k, \Delta P, R_D, s \text{ or } \rho$

The diameter of the pipe must be given or assumed. The fluids' velocities must be in the range of 3-10 ft/sec. The following assumes that the cold fluid is in the inner pipe.

(1) The heat duty Q, Btu/hr, is:

$$Q = WC(T_1 - T_2) = wc(t_2 - t_1)$$
(8-53)

(2) The log mean temperature difference (LMTD).

LMTD =
$$\frac{(T_1 - t_2) - (T_2 - t_1)}{\ln\left\{\frac{T_1 - t_2}{T_2 - t_1}\right\}}$$
(8-5)

Inner Pipe

(3) The flow area, a_p :

$$a_{p} = \frac{\pi D^{2}}{4}, \quad \text{ft}^{2}$$
(8-54)

(4) The mass velocity, G_p

$$G_{p} = \frac{w}{a_{p}}, \quad \frac{lb}{ft^{2}hr}$$
(8-55)

(5) The Reynolds number, Re_t

$$\operatorname{Re}_{t} = \frac{D.G_{p}}{\mu}$$
(8-56)

(6) The Prandtl number, Pr_t

$$\Pr_{t} = \frac{c.\mu}{k} \tag{8-57}$$

(7) The heat transfer coefficient, h_i , Btu/hrft² F

$$\frac{\mathbf{h}_{i} \cdot \mathbf{D}}{k} = \mathbf{C} \left(\frac{\mathbf{D} \cdot \mathbf{G}_{p}}{\mu}\right)^{0.8} \left(\frac{\mathbf{c} \cdot \mu}{k}\right)^{0.33} \left(\frac{\mu}{\mu_{w}}\right)^{0.14}$$
(8-58)

h_i can be expressed by

$$h_i = C.\left(\frac{k}{D}\right) Re_t^{0.8} Pr_t^{0.33}$$
 (8-59)

where $\mu/\mu_w = 1$

 $\ddot{C} = 0.021$ for gases C = 0.023 for non-viscous fluids C = 0.027 for viscous fluids

converting h_i to $h_{io},$ the heat transfer coefficient referred to the pipe outside diameter, $Btu/hrft^{2\circ}F$

$$h_{io} = h_{i} \left(\frac{\text{inside dia. of inner pipe}}{\text{outside dia. of inner pipe}} \right)$$
$$= h_{i} \cdot \frac{\text{ID}}{\text{OD}}$$
(8-60)

Annulus

(8) The flow area, a_a :

$$a_a = \frac{\pi (D_o^2 - D_i^2)}{4}, \quad \text{ft}^2$$
 (8-61)

(9) The equivalent diameter, D_e

$$D_{e} = \frac{4 \times \text{flow area}}{\text{wetted perimeter}}$$
$$= \frac{D_{e}^{2} - D_{i}^{2}}{D_{i}}, \quad \text{ft}$$
(8-62)

(10) The mass velocity, G_a

$$G_a = \frac{w}{a_a}, \quad \frac{lb}{ft^2 hr}$$
(8-63)

(11) The Reynolds number, Re_a

$$\operatorname{Re}_{a} = \frac{D_{e} \cdot G_{a}}{\mu}$$
(8-64)

(12) The Prandtl number, Pr_a

$$\Pr_a = \frac{c \cdot \mu}{k} \tag{8-65}$$

(13) The heat transfer coefficient, h_0

If $Re_a < 2100$

$$h_o = 1.86 \frac{k}{D_e} Re_a^{0.33} Pr_a^{0.33} \left(\frac{D_e}{L}\right)^{0.33}$$
 (8-66)

If $\text{Re}_{a} > 2100$

$$h_o = C. \frac{k}{D_e} Re_a^{0.8} Pr_a^{0.33}$$
 (8-67)

(14) The overall coefficient, U, Btu/hr. ft.² $^{\circ}F$

$$\frac{1}{U} = \frac{1}{h_{io}} + \frac{1}{h_o} + R_D + R_w$$
(8-68)

(15) The heat transfer area, A, ft^2

$$A = \frac{Q}{U_{\rm D}.\,\text{LMTD}} \tag{8-69}$$

Vapor Service

(16) The heat transfer coefficient, h_i, Btu/hrft²° F

$$h_{i} = \frac{0.0144C_{p}G^{0.8}}{D_{e}^{0.2}}$$
(8-70)

and

$$\frac{1}{h_{if}} = \frac{1}{h_i} + Fol \qquad (h_i \text{ is corrected for fouling})$$
(8-71)

Shell Side (Finned Tube)

Figure 8-13 shows examples of extended surfaces on one or both sides of heat exchangers.

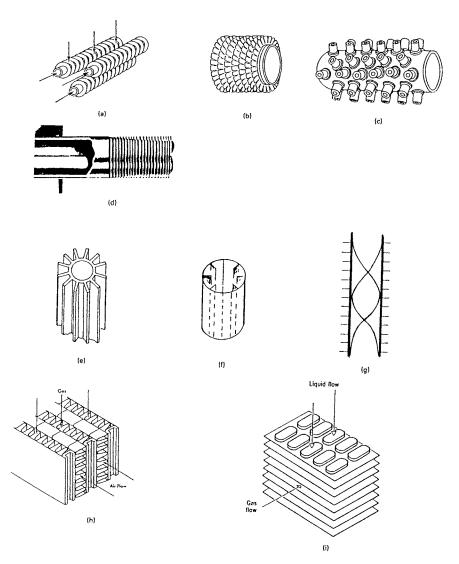


Figure 8-13. Examples of extended surfaces on one or both sides. (a) Radial fins. (b) Serrated radial fins. (c) Studded surface. (d) Joint between tubesheet and low fin tube with three times bare surface. (e) External axial fins. (f) Internal axial fins. (g) Finned surface with internal spiral to promote turbulence. (h) Plate fins on both sides. (i) Tubes and plate fins. Source: S. M. Walas, *Chemical Process Equipment: Selection and Design*, Butterworths Series in Chemical Engineering, 1988.

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(17) The equivalent diameter, D_e , ft

$$D_{e} = \frac{4NFA}{\pi(D_{s,i} + D_{t,o}) - N\theta + 2HN}$$
(8-72)

(18) The net free cross-sectional area, NFA, shell side, ft²

$$NFA = CSA - NH\theta$$
(8-73)

(19) The cross-sectional area without fins, CSA, ft^2

$$CSA = \frac{\pi}{4} \left(D_{s,i}^2 - D_{t,o}^2 \right)$$
(8-74)

(20) Finned surface area, ft²

$$A_{f} = 2HN \tag{8-75}$$

(21) Outside heat transfer area of tube, ft^2/ft (for finned tubes includes fin area)

$$A_{o} = \pi D_{t,o} + A_{f} \tag{8-76}$$

(22) Parameter for fin efficiency, X

$$X = H\left(\frac{h_{if}}{6K_{t}\theta}\right)^{0.5}$$
(8-77)

(23) The fin efficiency, e

$$e = \frac{\tanh X}{X} = \frac{1}{X} \left[\frac{\exp(X) - \exp(-X)}{\exp(X) + \exp(-X)} \right]$$
(8-78)

(24) The effective surface efficiency for fins, e_{eff} ,

$$e_{eff} = e\left(\frac{A_{f}}{A_{o}}\right) + \left(1 - \frac{A_{f}}{A_{o}}\right)$$
(8-79)

(25) The corrected film heat transfer coefficient, h_{ifd} , Btu/ft²hr°F $h_{ifd} = (h_{if})(e_{eff})$ (8-80)

(26) The overall heat transfer coefficient, U, Btu/ft²hr°F

Heat Transfer 635

$$\frac{1}{U} = \left(\frac{\Delta w}{K_{t}}\right)_{tube wall} + \frac{1}{h_{i}\left(\frac{A_{i}}{A_{o}}\right)_{tube}} + \frac{1}{(h_{ifd})_{shell}}$$
(8-81)

where

$$\frac{A_i}{A_o} = \frac{\pi D_{t,i}}{\pi D_{t,o} + 2HN}$$
(8-82)

Pressure Drop, $\Delta \mathbf{P}$

(27) The pressure drop in tubes is:

$$\Delta F = \frac{4 f G^2 L}{2 g \rho^2 D} \quad (ft \text{ of liquid})$$
(8-83)

where

$$f = \frac{16}{Re}$$
 for Re < 2,300 (8-84)

Using, Wilson et al. [15] correlation for commercial pipe

$$f = 0.0035 + \frac{0.264}{Re^{0.42}}$$
 for Re < 2,300 (8-85)

$$\Delta P_{p} = \left(\frac{\Delta F.\rho}{144}\right), \quad psi$$
(8-86)

Annulus

For an annulus, D'_e is:

$$D'_{e} = \frac{4\pi (D_{o}^{2} - D_{i}^{2})}{4\pi (D_{o} + D_{i})}$$

= D_{o} - D_{i} (8-87)

Reynolds number, Re_a

$$\operatorname{Re}_{a} = \frac{D'_{e}G_{a}}{\mu}$$
(8-88)

Pressure drop, ΔF_a , ft

$$\Delta F_{a} = \frac{4 f G_{a}^{2} L}{2 g \rho^{2} D_{e}^{\prime}}, \quad \text{ft}$$
(8-89)

The pressure drop due to the reversal of flow in the annulus for each hairpin is:

$$\Delta F_r = \frac{v^2}{2g}, \quad \text{ft/hair pin}$$
 (8-90)

The total annulus pressure drop is:

$$\Delta P_{a} = \left[\frac{(\Delta F_{a} + \Delta F_{r})\rho}{144}\right], \quad \text{psi}$$
(8-91)

Nomenclature

 $A = cross-sectional area, ft^2$

- A_i = inside heat transfer area-tube, ft²/ft
- $A_o =$ outside heat transfer area-tube, ft²/ft (for finned tubes includes fin area)

$$A_f = finned transfer area, ft^2$$

- $a_p = flow area, ft^2$
- \dot{C} = specific heat of hot fluid, Btu/lb°F
- c = specific heat of cold fluid, Btu/lb°F
- CSA = cross-sectional area, shell side without fins, ft²
 - D = inside diameter, ft
- D_i , D_o = for annuli, D_i is the outside diameter of inner pipe, D_o is the inside diameter of the outer pipe, inch
 - $D_{t,i}$ = tube inside diameter, inch
 - D_{to} = tube outside diameter, inch
 - D_{si} = shell inside diameter, inch
 - D_e = equivalent diameter for heat-transfer, ft
 - D_e = equivalent diameter for pressure drop, ft.
 - e = fin efficiency
 - e' = effective surface efficiency for fins
 - FOL = fouling factor, hrft²°F/Btu
 - f = friction factor, dimensionless
 - ΔF = pressure drop, ft

- ΔF_a = annulus pressure drop, ft
- ΔF_r = pressure drop due to reversal of flow in the annulus, ft
 - $G = mass velocity, lb/ft^2hr$
- $G_a = mass velocity in annulus, lb/ft^2hr$
- $G_p = mass velocity of inner pipe, lb/ft^2hr$
 - \dot{g} = acceleration due to gravity, 32.2 ft/sec² (4.18 × 10⁸ ft/hr²)
- H = fin height, inch
- h_i, h_o = heat transfer film coefficient for inside fluid and outside fluid, respectively, Btu/hr.ft.²⁰F
 - h_{io} = value of h_i when referred to the pipe outside diameter, Btu/hr. ft.²⁰F
 - h_{if} = heat transfer film coefficient corrected for fouling, Btu/hr.ft.²°F
 - h_{ifd} = corrected film heat transfer rate, Btu/hr.ft.²°F
 - ID = inside diameter, inch or ft
 - $j_{\rm H}$ = heat transfer factor, dimensionless
 - k = fluid thermal conductivity, Btu/hr.ft.²⁰F
 - K_t = thermal conductivity of tube material, Btu/hr.ft.²°F
 - L = pipe length, ft
- LMTD = log mean temperature difference, °F
 - NFA = net free cross-sectional area, shell side, ft^2
 - N = number of fins
 - OD = outside diameter, inch or ft
 - Pr = Prandtl number, dimensionless
 - $\Delta P = pressure drop, psi$
 - Q = exchanger duty, Btu/hr
 - $R_{\rm D}$ = fouling resistance, hrft²°F/Btu
 - Re = Reynolds number, dimensionless
 - $R_w =$ wall resistance
 - s = specific gravity
 - T_1 , T_2 = temperature of hot fluid, inlet and outlet respectively, °F
 - t_1, t_2 = temperature of cold fluid, inlet and outlet respectively, °F
 - \tilde{U} = overall heat transfer coefficient, Btu/hr.ft.²⁰F
 - V = fluid velocity, ft/sec
 - W = mass flow rate of hot fluid, lb/hr
 - w = mass flow rate of cold fluid, lb/hr
 - X = parameter used in fin efficiency
 - $\rho =$ fluid density, lb/ft³
 - μ , μ_w = fluid viscosity (flowing and at wall), cP
 - θ = fin thickness (normally 0.035 inch)
 - $\pi = 3.1415927$

Subscripts

a = annulusp = pipe

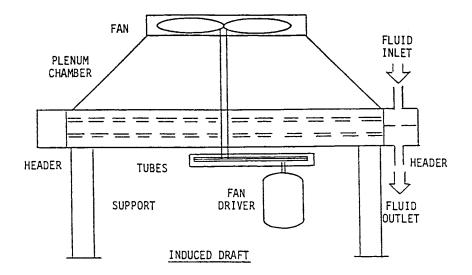
AIR COOLER DESIGN

Air coolers are widely used in the chemical process industries where there are shortages of cooling water and regulatory controls on thermal pollution. An economic comparison is often made between air and water cooling to determine which is advantageous. Air coolers can be used for cooling high pressure streams where the fluid is always inside the tubes. They minimize heat exchanger maintenance costs because descaling of the water-contact surface is eliminated. In addition, there is some savings in first cost because no additional surface area is required to allow for water side fouling between descaling. However, the exchangers are larger because the gas film coefficient is smaller. Air coolers eliminate problems of temperature rise and chemical pollution of water resources.

Air cooled heat exchangers have rectangular bundles containing several rows of tubes, horizontally aligned and vertically offset. Air flows vertically upward across the tube bank. The flow can be induced by fans above the bundle or forced by fans below the bundle. The heat transfer is countercurrent, because the hot fluid enters at the top of the bundle and flows downward through successive passes. The cost of an air cooler depends on the length of the tubes and the number of the tube rows.

A forced draft air cooler design has a power consumption advantage, if the temperature rise of the air is high. It also allows a more convenient and economical mounting arrangement when a number of bundles and services are to be combined in a single unit. In addition, it develops a high turbulence, and consequently, higher heat transfer coefficients. Conversely, an induced draft design provides a more even distribution of air across the bundles, and for a given bundle elevation, affords more space for location of additional plant equipment beneath the unit. Induced draft units are less likely to recirculate hot exhaust air, because the exit air velocity is two or three times that of a forced draft unit. These units give higher pressure drops and are more expensive. Figure 8-14 illustrates forced and induced air cooled exchanger arrangements.

Standard air coolers are in widths of 8, 10, 12, 16, or 20-ft lengths of 4-40ft, and stacks of 3–6 rows of tubes. Tubes are 0.75–1.0 inch OD



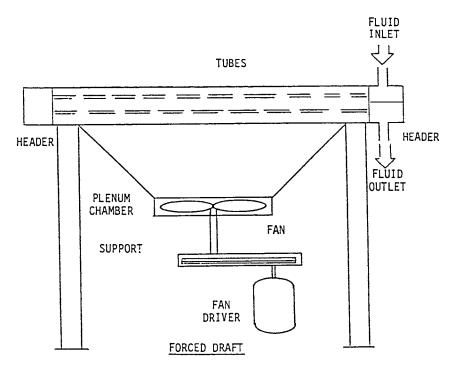


Figure 8-14. Forced and induced draft air cooled exchangers.

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with 7–11 fins/inch each 0.5–0.625 in height and provide a total surface 15–20 times greater than the bare surface of the tube. Fans are 4–12 ft/ diameter; developed pressures are 0.5–1.5 inch water, and require power inputs of 2–5 hp/MBtu/hr or about 7.5 hp/100 ft² of exchanger cross section. Spacings are 1.8 times the width of the cooler. Face velocities are about 10ft/sec at a depth of three rows and 8ft/sec at a depth of six rows. The rating methods for air cooled exchangers are outlined by Cook [16] and Ganapathy [17]. Here, the design of air coolers is based upon correlations, tables, and graphs presented by Smith [18] and Brown [19] and fitted to a series of equations developed by Blackwell [20].

The Equations

(1) The number of tube rows, R, is:

$$\mathbf{R} = \mathbf{C}_{1} + \mathbf{C}_{2} \ln \left[\frac{\mathbf{T}_{2} - \mathbf{t}_{2}}{\mathbf{U}} \right]$$
(8-92)

where $C_1 = 3.1679$ $C_2 = 3.7948$

(2) The area ratio is:

$$\frac{A}{FA} = C_3(R)^{C_4}$$
(8-93)

where $C_3 = 1.2557$ $C_4 = 1.0031$

(3) The face velocity of air, FV, ft/min.

$$FV = C_5(C_6)^R \tag{8-94}$$

where $C_5 = 720.8542$ $C_6 = 0.9530$

Typical face velocities (FVs) used for design are in Table 8-15. These values result in air cooled heat exchangers that approach an optimum cost [21]. This takes into account the purchase cost, the cost of installation, and the cost of power to drive the fans.

Table 8-16 lists an estimate of the outlet air temperature, based on 90° to 95° F design ambient air temperature.

(4) Estimated air outlet temperature, t_1 , $^{\circ}F$

Table 8-15 Design Face Velocities for Air-Cooled Exchangers

	FACE VELOCITY, ft/min (m/sec.)			
Number of tube rows	8 fins/in. (315 fins/m) 2.375 in. (0.0603 m) pitch	10 fins/in. (394 fins/m) 2.375 in. (0.0635 m) pitch	10 fins/in. (394 fins/m) 2.5 in. (0.0635 m) pitch	
3	650 (3.30)	625 (3.18)	700 (3.56)	
4	615 (3.12)	600 (3.05)	600 (3.35)	
5	585 (2.97)	575 (2.92)	625 (3.18)	
6	560 (2.84)	550 (2.79)	600 (3.05)	

Source: Chopey and Hicks [21].

Table 8-16 Estimated Outlet Air Temperature for Air-Cooled Exchangers

PROCESS INLET TEMPERATURE °C	OUTLET AIR TEMPERATURE °C		
	U=50	U=100	U=150
175	90	95	100
150	75	80	85
125	70	75	80
100	60	65	70
90	55	60	65
80	50	55	60
70	48	50	55
60	45	48	50
50	40	41	42

Source: Chopey and Hicks [21].

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$$t_1 = 0.005 U \left[\left(\frac{T_1 + T_2}{2} \right) - t_2 \right] + t_2$$
 (8-95)

(5) The effective log mean temperature difference, °F

LMTD =
$$\frac{(T_1 - t_2) - (T_2 - t_1)}{\ln\left\{\frac{T_1 - t_2}{T_2 - t_1}\right\}}$$
(8-5)

(6) The heat duty Q, Btu/hr, is:

$$Q = wc\Delta t \tag{8-96}$$

where w =fluid flow rate, lb/hr

c = specific heat capacity, Btu/lb°F Δt = temperature difference, °F

(7) The bare tube surface area based on the tube, OD, ft^2

$$A = \frac{Q}{(U)(LMTD)}$$
(8-97)

(8) The face area of bundle FA, ft^2

$$FA = \frac{A}{C_3(R)^{C_4}}$$
(8-98)

(9) The calculated outlet air temperature, t'_1 , $^{\circ}F$

$$t_1' = \frac{Q}{(1.08)(FA)(FV)} + t_2$$
(8-99)

(10) The air flow over tubes F, std.ft³/min

$$F = (FA)(FV)$$
 (8-100)

(11) The ratio of the bare tube surface area based on the tube OD to the fan horsepower

$$\frac{A}{Bhp} = C_7 + C_8(R)$$
(8-101)
where $C_7 = 7.4212$
 $C_8 = 12.5342$

(12) The ratio of air cooler weight to the face area of bundle is:

$$\frac{Wt}{FA} = C_9 + C_{10}(R)$$
(8-102)

where $C_9 = 36.4$ $C_{10} = 9.35$

(13) The tube bundle width, W, ft

$$W = \frac{FA}{L}$$
(8-103)

(14) The area available, A_v , ft^2

$$A_v = \pi . N_r . N_r . D_o . L, ft^2$$
 (8-104)

where $N_r =$ number of rows

 $\dot{N_t}$ = number of tubes per row = (tube bundle width)/(tube spacing) = W/s L = tube length, ft D_o = tube outside diameter, ft

If the area available, A_v , is less than the area required, A, increase the bundle width, and calculate the new outlet air temperature. Re-calculate LMTD, the new required area, and the new available area.

(15) The air-side heat transfer coefficient, h_a , Btu/hrft²°F. The air side coefficient is determined on the basis of the outside surface of a bare tube. h_a is expressed as:

$$h_a = 8(FV)^{0.5}$$
 for 10 fins per inch (8-105)

$$h_a = 6.75 (FV)^{0.5}$$
 for 8 fins per inch (8-106)

(16) The tube wall heat transfer coefficient h_w , Btu/hrft²°F.

$$h_{w} = \frac{2.k}{D_{o} - D_{i}}$$
(8-107)

where k = thermal conductivity, Btu/hrft°F

 $D_0 =$ outside diameter of tube, ft

 $D_i = inside diameter of tube, ft$

(17) The overall heat transfer coefficient U, Btu/hrft²°F is

$$\frac{1}{U} = \frac{1}{h_a} + \frac{1}{h_i(D_i/D_o)} + \frac{1}{h_w} + \frac{1}{h_s}$$
(8-108)

where $h_i = inside$ of tube heat transfer coefficient $h_s = fouling coefficient$

The Air-Side Pressure Drop, ΔP_a (inch H₂O)

The following equations are used to calculate the pressure drop on the air side [21].

 $\Delta P_a = 0.0047 N_r (FV/100)^{1.8}$ for 10 fins per inch, 2.375-inch spacing (8-109)

 $\Delta P_a = 0.0044 N_r (FV/100)^{1.8}$ for 8 fins per inch, 2.375-inch spacing (8-110)

 $\Delta P_a = 0.0037 N_r (FV/100)^{1.8}$ for 10 fins per inch, 2.5 inch spacing (8-111)

where N_r = the number of tube rows FV = face velocity, ft/min ΔP_a = inch H₂O

Nomenclature

A = bare-tube surface area, ft^2

 $A_v = available area, ft^2$

Bhp = fan horsepower

c = specific heat of fluid, Btu/lb°F

 $C_1, C_2, C_3 = constants$

 $C_4, C_5, C_6 = constants$

 $C_7, C_8, C_9 = constants$

 $C_{10} = constant$

 D_i = inside diameter of tube, ft

 D_0 = outside diameter of tube, ft

 $F = air flow over tubes, stdft^3/min$

 $FA = face area of bundle, ft^2$

FV = face velocity of air, ft/min

 h_a = air-side heat transfer coefficient, Btu/hrft²°F

 h_i = inside of tube heat transfer coefficient, Btu/hrft²⁰F

 $h_s =$ fouling coefficient, Btu/hrft²°F

 h_w = tube wall heat transfer coefficient, Btu/hrft² F

k = thermal conductivity, Btu/hrft°F

L = pipe length, ftLMTD = log mean temperature difference, $^{\circ}F$ $N_r =$ number of rows N_t = number of tubes per row $\Delta P_a = air-side pressure drop, inch H_2O$ Q = exchanger duty, Btu/hrR = number of tube rows T_1 = outlet process fluid temperature, °F T_2 = inlet process fluid temperature, °F t_1 = estimated air outlet temperature, °F t_1 = calculated outlet air temperature, °F $t_2 = inlet air temperature, °F$ $\Delta t = temperature difference, ^{\circ}F$ U = overall heat transfer coefficient (based on bare-tube O.D.),Btu/hrft²⁰F W = tube bundle width, ftw =fluid flow rate, lb/hr Wt = air cooler weight, lb $\pi = 3.1415926$

HEAT TRACER REQUIREMENTS FOR PIPELINES AND HEAT LOSS FROM INSULATED PIPELINES

Pipelines conveying viscous fluids are maintained at an elevated temperature by means of heat tracing. Pipelines containing vapors may also be heat traced to prevent components from condensing out. The heat loss from pipes is often reduced by thermal insulation. The thickness of insulation depends on an economic analysis involving both capital cost and the cost of heat loss from the insulated line. Water lines are commonly insulated to avoid freezing. However, heat is inevitably lost from insulated lines, and if the cooling resulting from this heat loss cannot be tolerated, heat tracing the lines becomes necessary. There are cases where blockages occur in pipelines, which cannot be unblocked by flushing with a solvent or by blowing in steam or air, unless they are heat traced.

Many types of fluid heating media are used for heat tracing, such as steam, hot oil, or Dowtherm. Electric heating tape may be used. Steam tracing is selected for about 60 percent of pipe footage in chemical process plants Lam and Sandberg [22] have provided a comprehensive checklist and criteria between steam and electric tracing. Table 8-17 lists a typical checklist for a petroleum refinery plant.

Table 8-17 Checklist to Decide Between Steam and Electric Heat Tracing

	Electric	Steam
PRE-EXISTING PROJECT CONSTRAINTS		
Heat tracing recommended in company (Yes/no)	Yes	Yes
engineering practices		
Plant preference		Yes
PROJECT DESIGN PHASE CRITERIA		
Total feet of traced pipe	4,700	4,700
Cost of electricity (\$/kWh) and steam energy (\$/1,000 lb)	\$0.067	Consider free
Estimated annual maintenance cost per foot of tracing (\$/ft.)	\$1.67	\$1.33
Number of heat-tracing circuits required	50	65
Needed accuracy of temperature control (degrees)	±50°	±50°
Temperature control cost per circuit for needed accuracy (\$)	No extra	No extra
Cost of monitoring one circuit in distributed control system (\$)	NA*	NA*
Design time per circuit (h)	0.5	1.5
Design tools available for engineers (Good, Fair, Poor, None)	Good	Good
Capital cost of steam capacity, condensate return, etc. (\$)		Low
Capital cost of required electrical power (\$)	Low	
Other		
PROJECT INSTALLATION CRITERIA		
Labor cost to install tracing and accessories (\$/ft)	\$30.00	\$40.00
Training time per plant laborer (h)	20	20
Labor and overhead costs (\$/h)	\$23.50	\$23.50
Total installed heat-tracing costs (\$/ft)	\$62.00	\$74.00
Other		
PROJECT OPERATION CRITERIA		
Annual maintenance required (h/ft.)	0.03	0.05
Annual cost for replacement parts (\$/ft.)	\$1.00	\$0.80
Total annual maintenance cost (\$)	\$6,182	\$6,807
Total annual operating cost (\$) Other	\$7,179	\$6,807

*NA = not applicable, as the plant does not have a distributed control system capable of monitoring heat tracing

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The length of a traced line varies from a few feet in a process area to a few thousand feet between offsites and the process area, as well as to hundreds of miles involving underground pipelines conveying crude oil. The most common steam tracing lines are 3/8-inch and 1/2-inch o.d. copper or stainless steel tubing. Larger tubing of 5/8-inch and 3/4-inch o.d. can be used, but is more expensive. The 3/8-inch tube is more easily plugged by debris or sediment, and is therefore used less than 1/2-inch tubing. Copper is preferred for its heat transfer characteristics, but stainless steel generally provides better resistance in corrosive situations. The length of a single trace tube (for example, from steam supply valve to steam trap) is limited by the pressure drop in the trace. In addition, the trap should have a condensate drainage capacity to match the heating load. At steam pressure of 100 psig or higher, Kohli [23] recommends that the length of a single trace should not exceed 200 ft. Conversely, if the steam pressure is lower, a tracer length of 100 ft is recommended. The trace tube is wired to the pipeline to provide greater contact. The conductive heat transfer rate is often low, but this may be increased by putting a layer of heat conducting cement (graphite mixed with sodium silicate or other binders) between the trace and the pipeline. This results in better surface for conductive heat transfer. Figure 8-15 shows various configurations of tracer and lagging.

The following equations will allow the design engineer to calculate the heat loss and tracing requirements for any given pipeline, using any hot fluid medium as the heat source. The equations will calculate:

- The surface temperature of insulated traced pipe.
- Total heat transferred per 100ft of pipe.
- Total heat transferred for the entire pipeline.
- Flow rate of hot media.
- Total number of heat tracers required with and without transfer cement.

Kern [3] and others have shown that the heat transferred through an insulated pipe involves four resistances:

- 1. The film resistance on the inside wall of the pipe.
- 2. Heat resistance through the pipewall.
- 3. Heat resistance through the insulation.
- 4. Air film resistance on the outside of the insulation.

The first two resistances are negligible, and therefore are neglected.

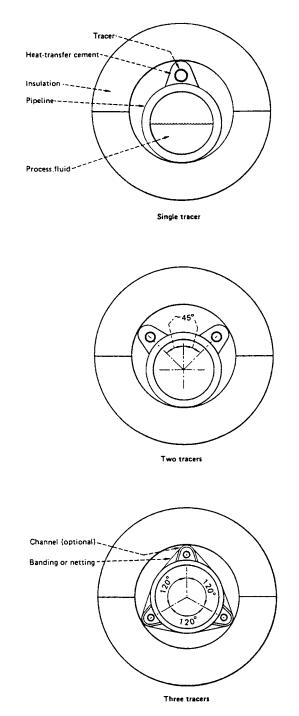


Figure 8-15. Configuration of heat tracers and lagging.

The Equations

(1) The average temperature of the hot medium, °F

$$T_{m,avg} = 0.5(T_{mi} + T_{mo})$$
(8-112)

(2) The average temperature of pipe and tracer, $^\circ F$

$$T_{avg} = 0.5(T_{m,avg} + T_p)$$
(8-113)

(3) The inside diameter of insulation, inch

$$D_i = OD + TAL$$
(8-114)

(4) The outside diameter of insulation, inch

$$D_{o} = D_{i} + 2.T_{k}$$
 (8-115)

(5) Heat lost per foot of pipe Q, Btu/hr.ft

$$Q = \frac{2\pi K_{i}(T_{a} - T_{s})}{\ln\left(\frac{D_{o}}{D_{i}}\right)}$$
(8-116)

The heat lost per foot of pipe can be expressed in terms of the film heat transfer coefficient to air corrected for wind, Btu/hrft²°F

$$Q = \frac{h_a \pi D_o (T_s - T_{air})}{12}$$
(8-117)

Using Equations 8-116 and 8-117 and rearranging the terms gives

$$\frac{2\pi K_{i}(T_{a} - T_{s})}{\ln\left(\frac{D_{o}}{D_{i}}\right)} = \frac{h_{a}\pi D_{o}(T_{s} - T_{air})}{12}$$
(8-118)

That is,

$$T_a - T_s = C(T_s - T_{air})$$
 (8-119)

where

$$C = (h_a) \left(\frac{D_o}{12} \right) \left(\frac{1}{2K_i} \right) \left(\ln \frac{D_o}{D_i} \right)$$
(8-120)

Equation 8-119 can be expressed in terms of T_s as follows:

$$T_{s}(l+C) = T_{a} + CT_{air}$$
 (8-121)

and

$$T_s = \frac{T_a + CT_{air}}{1 + C}$$
(8-122)

Equations 8-120 and 8-122 involve two unknowns, h_a and T_s . An initial value of h_a is assumed and T_s is calculated. McAdams [24] indicates that h_a usually varies from 1.64 to 5.30 for pipes in still air, depending upon pipe size, surface, and air temperature. Table 8-18 gives values of h_a for pipes in still air, and Table 8-19 lists the correction factor for h_a at different wind velocities.

(6) The combined convective and radiative heat transfer coefficient $(h_c + h_r)$ is given by

$$h_{c} + h_{r} = \frac{564}{D_{o}^{0.19} [273 - (T_{s} - T_{air})]}$$
(8-123)

(7) The wind factor, WF:

WF=
$$C_0 + C_1 (T_s - T_{air}) + C_2 (T_s - T_{air})^2$$
 (8-124)

	(-;	s •a/ ·	(1.01.01	, annagg	jou pipe	•s - •w/		
Nominal pipe dia., inch.	50	100	150	200	250	300	400	500
1/2	2.12	2.48	2.76	3.10	3.41	3.75	4.47	5.30
1	2.03	2.38	2.65	2.98	3.29	3.62	4.33	5.16
2	1.93	2.27	2.52	2.85	3.14	3.47	4.18	4.99
4	1.84	2.16	2.41	2.75	3.01	3.33	4.02	4.83
8	1.76	2.06	2.29	2.60	2.89	3.20	3.83	4.68
12	1.71	2.01	2.24	2.54	2.82	3.12	3.83	4.61
24	1.64	1.93	2.15	2.45	2.72	3.03	3.70	4.48

Table 8-18Values of h_a for Pipes in Still Air $(t_s - t_a) \ ^\circ F$ (For an unlagged pipe $t_s = t_w$)

Source: McAdams [24].

Wind velocity, m/hr	100	200	300	400	500
2.5	1.46	1.43	1.40	1.36	1.32
5.0	1.74	1.69	1.64	1.59	1.53
10.0	2.16	2.10	2.02	1.93	1.84
15.0	2.50	2.42	2.33	2.27	2.08
20.0	2.76	2.69	2.58	2.45	2.30
25.0	2.98	2.89	2.78	2.64	2.49
30.0	3.15	3.06	2.94	2.81	2.66
35.0	3.30	3.21	3.10	2.97	2.81

Table 8-19Correction Factor for h_a at Different Wind Velocities($t_s - t_a$), °F (For an unlagged pipe, $t_S = t_W$)

Source: Kohli [23]

where $C_0 = 2.814$ $C_1 = -0.00038857$ $C_2 = -0.0000012857$

For zero wind condition,

$$C_0 = 1.0$$

$$C_1 = 0.0$$

 $C_2 = 0.0$

(8) The new film heat transfer coefficient to air corrected for wind, $h_a(Btu/hrft^{2\circ} F)$

$$h_a = (h_c + h_r)(WF)$$
 (8-125)

(9) The total heat lost from pipeline, Q_t , Btu/hr

$$Q_t = (Q)(L_{\text{TOTAL}}) \tag{8-126}$$

where L_{TOTAL} = total length of pipe, ft

(10) The flow rate of hot medium W, lb/hr

$$W = \frac{Q_{t}}{C_{p}(T_{mi} - T_{mo})}$$
(8-127)

(11) The number of tracers required without heat transfer cement is

$$N_{TWOC} = \frac{Q}{a(T_{m,avg} - T_p)}$$
 (8-128)

where a = thermal conductance tracer to pipe without cement, Btu/hr°F(ft of pipe)

(12) The number of tracers required with heat transfer cement:

$$N_{\rm TWC} = \frac{Q}{b(T_{\rm m,avg} - T_{\rm p})}$$
(8-129)

where b = thermal conductance tracer to pipe with cement, Btu/hr°F(ft. of pipe)

Table 8-20 lists the values of a and b.

Blackwell [20] recommends approximately 1 1/4 inch between pipe and insulation to accommodate a 1/2-inch tracer line and heat transfer cement. He further recommends twice this value for three or more tracers. A 3/8 inch to 1 inch of space may be required for smaller tracers. Tracers are normally spaced equidistantly around the pipe, and are run parallel to it. In the case of heat lost from an insulated pipeline without tracing, the hot medium inlet and outlet temperatures are set to equal the temperature of the pipe.

Nomenclature

- a = thermal conductance, tracer to pipe without heat transfer cement, Btu/hr°F(ft of pipe)
- b = thermal conductance, tracer to pipe with cement, Btu/hr°F(ft of pipe)

 C_0, C_1, C_2 = constants for wind factor equation

Table 8-20Thermal Conductance Tracer to Pipe

Tube size, inch	а	b
3/8	0.295	3.44
1/2	0.393	4.58
5/8	0.490	5.73

Source: Kohli [23].

 C_{p} = specific heat of hot medium, Btu/lb°F

- $\dot{\mathbf{D}}_{i}$ = inside diameter of insulation, inch
- D_0 = outside diameter of insulation, inch
- $h_a = film$ heat transfer coefficient to air (corrected for wind), (Btu/hrft²°F)

 $h_c + h_r =$ combined convective and radiative heat transfer coefficients, (Btu/hrft²°F)

 K_i = thermal conductivity of insulation, Btu/hrft²(°F/ft)

 L_{TOTAL} = total pipeline length, ft

 N_{TWC} = number of tracers required with heat transfer cement

 N_{TWOC} = number of tracers required without heat transfer cement

Q = heat lost per ft of pipe, Btu/hr ft

 Q_t = total heat lost from pipeline, Btu/hr

 T_a = average temperature of pipe and tracer, °F

 $T_{air} = air temperature, ^{\circ}F$

TAL = allowance for tracer diameter, inch

 T_{maxy} = average temperature of hot medium, °F

 T_{mi} = inlet temperature of hot medium, °F

 T_{mo} = outlet temperature of hot medium, °F

 T_{n} = temperature in pipe, °F

- \vec{T}_s = outside surface temperature of insulation, °F
- W =flow rate of hot medium, lb/hr

 $W_F = wind factor$

BATCH HEATING AND COOLING OF FLUIDS

Heating or cooling of process fluids in a batch operated vessel is common in the chemical process industries. The process is an unsteady state in nature, because the heat flow, the temperature, or both vary with time at a fixed point. The time required for the heat transfer can be modified by increasing the agitation of the batch fluid, the rate of circulation of the heat transfer medium in a jacket or coil or both or the heat transfer area. Bondy and Lippa [25] have compiled a collection of correlations of heat transfer coefficients in agitated vessels. Batch processes are sometimes disadvantageous because:

- Use of the heating or cooling medium is intermittent.
- The liquid being processed is not readily available.
- The requirements for treating time requires holdup.
- Cleaning or regeneration is an integral part of the total operating period.

The variables in batch heating or cooling processes are surface requirement, time, and temperature. Heating a batch may be by external means, such as a jacket or coil, or by withdrawing and recirculating it through an external heat exchanger. In either case, assumptions are made to facilitate calculation:

- 1. The overall heat transfer coefficient, U, is constant for the process and over the entire surface.
- 2. Liquid flow rates are at steady state.
- 3. Specific heats are constant for the process.
- 4. The heating or cooling medium has a constant inlet temperature.
- 5. Agitation gives a uniform batch fluid temperature.
- 6. There is no phase change.
- 7. Heat losses are negligible.

Batch Heating: Internal Coil, Isothermal Heating Medium

When an agitated batch containing M of fluid with specific heat, c, and initial temperature, t, is heated with an isothermal condensing heating medium, T_1 , the batch temperature t_2 , at any time, θ , can be derived by the differential heat balance. For an unsteady state operation as shown in Figure 8-16, the total number of heat transferred is q['], and per unit time, θ , is:

I II III IV

$$dq = \frac{dq'}{d\theta} = Mc \frac{dt}{d\theta} = UA\Delta t$$
 (8-130)
Accumu- Transfer
lation in rate
the batch
 $\Delta t = T_1 - t$ (8-131)

Equating III and IV gives

$$Mc \frac{dt}{d\theta} = UA\Delta t$$
(8-132)

Rearranging Equation 8-132 gives:

$$\frac{\mathrm{dt}}{\mathrm{\Delta t}} = \frac{\mathrm{UA}}{\mathrm{Mc}} \,.\,\mathrm{d\theta} \tag{8-133}$$

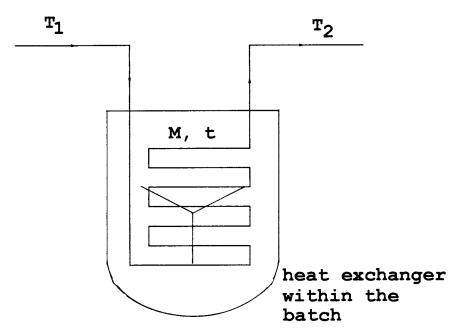


Figure 8-16. Agitated batch vessel.

$$\int_{t_1}^{t_2} \frac{\mathrm{d}t}{\mathrm{T}_1 - \mathrm{t}} = \frac{\mathrm{UA}}{\mathrm{Mc}} \int_0^{\theta} \mathrm{d}\theta \qquad (8-134)$$

Integrating Equation 8-134 from t_1 to t_2 while the time passes from 0 to θ yields:

$$\ln\left(\frac{T_1 - t_1}{T_1 - t_2}\right) = \frac{UA}{Mc} \cdot \theta$$
(8-135)

where A = heat transfer surface

- c = specific heat of liquid
- M = weight of batch liquid

 T_1 = heating medium temperature

- t_1 = initial batch temperature
- $t_2 =$ final batch temperature U = overall heat transfer coefficient
- $\theta = time$

Batch Cooling : Internal Coil Isothermal Cooling Medium

Consider the same arrangement containing M of liquid with specific heat, c, and initial temperature, T_1 , cooled by an isothermal vaporizing medium of temperature, t_1 . If T is the batch temperature at any time, θ , then

$$\frac{\mathrm{d}\mathbf{q}'}{\mathrm{d}\theta} = -\mathrm{Mc}\,\frac{\mathrm{d}\mathrm{T}}{\mathrm{d}\theta} = \mathrm{U}\mathrm{A}\Delta\mathrm{t} \tag{8-136}$$

where

 $\Delta t = T - t_1 \tag{8-137}$

Then

$$-Mc \frac{dT}{d\theta} = UA\Delta t$$
 (8-138)

Substituting Equation 8-137 into Equation 8-138 and rearranging gives:

$$-\int_{T_1}^{T_2} \frac{dT}{T-t_1} = \int_0^{\theta} \frac{UA}{Mc} \,.\,d\theta$$
 (8-139)

Integrating from T_1 to T_2 while the time passes from 0 to θ gives:

$$\ln\left(\frac{T_1 - t_1}{T_2 - t_1}\right) = \frac{UA}{Mc} \cdot \theta$$
(8-140)

where A = heat transfer surface

c = specific heat of liquid

- M = weight of batch liquid
- T_1 = initial batch temperature
- $T_2 =$ final batch temperature
 - $t_i = cooling medium temperature$
- U = overall heat transfer coefficient
- $\theta = time$

Batch Heating with Non-Isothermal Heating Medium

The non-isothermal heating medium has a constant flow rate, W_h , specific heat, C_h , and inlet temperature, T_l , but a variable outlet temperature. For an unsteady state operation,

I II III IV

$$\frac{dq'}{d\theta} = Mc \frac{dt}{d\theta} = W_h C_h (T_1 - T_2) = UA\Delta t_{LMTD}$$
(8-141)

The log mean temperature difference, Δt_{LMTD} is:

$$\Delta t_{\rm LMTD} = \frac{T_1 - T_2}{\ln\left(\frac{T_1 - t}{T_2 - t}\right)}$$
(8-142)

Equating III and IV in Equation 8-141 and rearranging gives:

$$\frac{W_{h}C_{h}(T_{1}-T_{2})}{UA} = \frac{T_{1}-T_{2}}{\ln\left(\frac{T_{1}-t}{T_{2}-t}\right)}$$
(8-143)

Equation 8-143 becomes:

$$\ln\left(\frac{T_1 - t}{T_2 - t}\right) = \frac{UA}{W_h C_h}$$
(8-144)

$$\frac{T_{1} - t}{T_{2} - t} = e^{UA/W_{h}C_{h}}$$
(8-145)

Rearranging Equation 8-145 gives:

$$T_{2} = t + \frac{T_{1} - t}{e^{UA/W_{h}C_{h}}}$$
(8-146)

where

$$\mathbf{K}_{\mathrm{I}} = \mathbf{e}^{\mathrm{U}\mathrm{A}/\mathrm{W}_{\mathrm{h}}\mathrm{C}_{\mathrm{h}}} \tag{8-147}$$

Equating II and III in Equation 8-141 and substituting Equation 8-146 in Equation 8-141 gives:

$$Mc \frac{dt}{d\theta} = W_{h}C_{h}\left(T_{1} - \left\{t + \frac{T_{1} - t}{K_{1}}\right\}\right)$$
(8-148)

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$$= W_{h}C_{h}\left(\frac{K_{1}-1}{K_{1}}\right)(T_{1}-t)$$
 (8-149)

$$\int_{t_{1}}^{t_{2}} \frac{dt}{T_{1} - t} = \int_{0}^{\theta} \frac{W_{h}C_{h}}{Mc} \left(\frac{K_{1} - 1}{K_{1}}\right) d\theta$$
(8-150)

Rearranging and integrating Equation 8-150 from t_1 to t_2 while the time passes from 0 to θ gives:

$$\ln\left(\frac{T_1 - t_1}{T_1 - t_2}\right) = \left(\frac{W_h C_h}{Mc}\right) \left(\frac{K_t - 1}{K_t}\right) \theta$$
(8-151)

where A = heat transfer surface

c = specific heat of batch liquid C_h = heating medium specific heat M = weight of batch liquid T₁ = heating medium temperature t₁ = initial batch temperature t₂ = final batch temperature U = overall heat transfer coefficient W_h = heating medium flow rate θ = time

Batch Cooling: Non-Isothermal Cooling Medium

When cooling a batch with internal coil and a non-isothermal cooling medium, the following equation can be applied.

$$\frac{\mathrm{d}\mathbf{q}'}{\mathrm{d}\theta} = -\mathrm{Mc}\frac{\mathrm{d}\mathbf{T}}{\mathrm{d}\theta} = \mathrm{W}_{\mathrm{c}}\mathrm{C}_{\mathrm{c}}(\mathrm{t}_{2} - \mathrm{t}_{1}) = \mathrm{U}\mathrm{A}\Delta\mathrm{t}_{\mathrm{LMTD}}$$
(8-152)

where $K_2 = e^{UA/W_cC_c}$

and

$$\ln\left(\frac{T_1 - t_1}{T_2 - t_1}\right) = \left(\frac{W_c C_c}{Mc}\right) \left(\frac{K_2 - 1}{K_2}\right) \theta$$
(8-153)

where A = heat transfer surface c = specific heat of liquid $C_c = coolant specific heat$

M = weight of batch liquid

 T_{I} = initial batch temperature

 T_2 = final batch temperature

 t_1 = initial coolant temperature

U = overall heat transfer coefficient

 $W_c = coolant flow rate$

 $\theta = time$

Batch Heating Through An External Heat Exchanger, Isothermal Heating Medium

Figure 8-17 illustrates the arrangement in which the fluid in the tank is heated by an external heat exchanger. The heating medium is isothermal, therefore any type of exchanger with steam in the shell and tubes can be used. The variable temperature out of the exchanger, t´,will differ from the variable tank temperature, t. An energy balance around the tank and the heat exchanger gives:

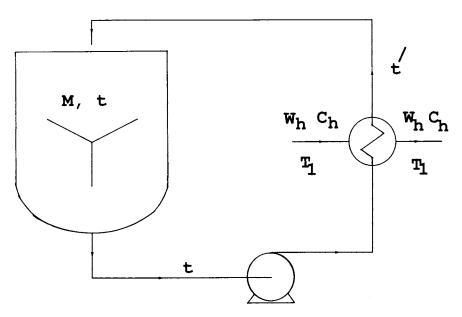


Figure 8-17. Batch heating through an external heat exchanger, isothermal heating medium.

$$I \qquad II \qquad III \qquad III$$

$$\frac{dq'}{dt} = Mc \frac{dt}{d\theta} = W_h C_h (t' - t) = UA\Delta t_{LMTD} \qquad (8-154)$$
Heat Heat enter- Transfer accumu- ing the rate in the lation in batch by external the batch recirculation exchanger

The log mean temperature difference, Δt_{LMTD} is:

$$\Delta t_{\rm LMTD} = \frac{(T_1 - t_1) - (T_1 - t')}{\ln\left(\frac{T_1 - t_1}{T_1 - t'}\right)}$$
(8-155)

$$=\frac{t'-t_{1}}{\ln\left(\frac{T_{1}-t_{1}}{T_{1}-t'}\right)}$$
(8-156)

Equating II and III in Equation 8-154 gives:

$$W_{h}C_{h}(t'-t) = UA\Delta t_{LMTD}$$
(8-157)

That is:

$$W_{h}C_{h}(t'-t) = UA \frac{(t'-t)}{\ln\left\{\frac{T_{1}-t_{1}}{T_{1}-t'}\right\}}$$
(8-158)

Rearranging Equation 8-158 gives

$$\ln\left(\frac{T_{1}-t}{T_{1}-t'}\right) = \frac{UA}{W_{h}C_{h}}$$
(8-159)

Equation 8-159 can be expressed as:

$$T_{1} - t = e^{UA/W_{h}C_{h}}(T_{1} - t')$$
(8-160)

where

$$K_3 = e^{UA/W_b C_b}$$

 $T_1 - t = K_3(T_1 - t')$
(8-161)

Therefore

$$t' = T_1 - \left(\frac{T_1 - t}{K_3}\right)$$
 (8-162)

Equating I and II in Equation 8-154

$$Mc \frac{dt}{d\theta} = W_{h}C_{h}(t'-t)$$
(8-163)

Substituting Equation 8-162 into Equation 8-163 and rearranging gives:

$$\frac{Mc}{W_{h}C_{h}} \cdot \frac{dt}{d\theta} = \left\{ T_{1} - \left(\frac{T_{1} - t}{K_{3}}\right) \right\} - t$$
(8-164)

$$=\frac{(K_3 - 1)(T_1 - t)}{K_3}$$
(8-165)

$$\int_{t_1}^{t_2} \frac{dt}{T_1 - t} = \left(\frac{K_3 - 1}{K_3}\right) \left(\frac{W_h C_h}{Mc}\right) \int_0^{\theta} d\theta$$
(8-166)

Rearranging and integrating Equation 8-166 from t_1 to t_2 while the time passes from 0 to θ gives:

$$\ln\left(\frac{\mathbf{T}_{1}-\mathbf{t}_{1}}{\mathbf{T}_{1}-\mathbf{t}_{2}}\right) = \left(\frac{\mathbf{K}_{3}-1}{\mathbf{K}_{3}}\right)\left(\frac{\mathbf{W}_{c}\mathbf{C}_{h}}{\mathbf{M}c}\right)\boldsymbol{\theta}$$
(8-167)

where A = heat transfer surface

- c = specific heat of batch liquid
- C_h = heating medium specific heat
- M = weight of batch liquid
- T_1 = heating medium temperature
 - t_1 = initial batch temperature

 t_2 = final batch temperature U = overall heat transfer coefficient W_h = heating medium flow rate θ = time

Batch Cooling Through an External Heat Exchanger, Isothermal Cooling Medium

When cooling a batch with an external heat exchanger and an isothermal cooling medium, the equation is:

$$\ln\left(\frac{T_1 - t_1}{T_2 - t_1}\right) = \left(\frac{W_c C_c}{Mc}\right) \left(\frac{K_4 - 1}{K_4}\right) \theta$$
(8-168)

where $K_4 = e^{UA/W_cC_c}$ A = heat transfer surface c = specific heat of liquid $C_c =$ coolant specific heat M = weight of batch liquid $T_1 =$ initial batch temperature $T_2 =$ final batch temperature $t_1 =$ initial coolant temperature U = overall heat transfer coefficient $W_c =$ coolant flow rate $\theta =$ time

Batch Cooling: External Heat Exchanger (Countercurrent Flow), Non-isothermal Cooling Medium

When cooling a batch with an external heat exchanger and a nonisothermal cooling medium, the following equation can be used.

$$\ln\left(\frac{T_1 - t_1}{T_2 - t_1}\right) = \left(\frac{K_5 - 1}{M}\right) \left(\frac{W_b W_c C_c}{K_5 W_c C_c - W_b c}\right) \theta$$
(8-169)

where $K_5 = exp\{UA(1/W_bc - 1/W_cC_c)\}$ A = heat transfer surface c = specific heat of liquid $C_c = coolant specific heat$ M = weight of batch liquid T_1 = initial batch temperature

 $T_2 =$ final batch temperature

 $t_i = initial \ coolant \ temperature$

U = overall heat transfer coefficient

 $W_{b} = batch flow rate$

 $W_c = coolant$ flow rate

 θ = time

Batch Heating: External Heat Exchanger and Non-isothermal Heating Medium

When heating a batch with an external heat exchanger and nonisothermal heating, the following equation applies:

$$\ln\left(\frac{\mathbf{T}_{1}-\mathbf{t}_{1}}{\mathbf{T}_{1}-\mathbf{t}_{2}}\right) = \left(\frac{\mathbf{K}_{6}-1}{\mathbf{M}}\right)\left(\frac{\mathbf{W}_{b}\mathbf{W}_{h}\mathbf{C}_{h}}{\mathbf{K}_{6}\mathbf{W}_{h}\mathbf{C}_{h}-\mathbf{W}_{b}\mathbf{c}}\right)\boldsymbol{\Theta}$$
(8-170)

where $K_6 = \exp\{UA(1/W_bc - 1/W_hC_h)\}$

A = heat transfer surface

c = specific heat of batch liquid

 C_h = heating medium specific heat

M = weight of batch liquid

 T_1 = heating medium temperature

 $t_i = initial batch temperature$

 t_2 = final batch temperature

 \overline{U} = overall heat transfer coefficient

 W_{b} = batch flow rate

 W_{h} = heating medium flow rate

 $\theta = time$

Batch Heating: External Heat Exchanger (1-2 Multipass Heat Exchangers), Non-isothermal Heating Medium

The procedure used for batch heating with external 1-2 multipass heat exchangers with non-isothermal heating media involves using the same heat balance as defined by the following equation:

$$\frac{\mathrm{d}\mathbf{q}'}{\mathrm{d}\theta} = \mathrm{Mc}\frac{\mathrm{d}\mathbf{t}}{\mathrm{d}\theta} = \mathrm{W}_{\mathrm{b}}\mathrm{c}(\mathbf{t}'-\mathbf{t}) = \mathrm{W}_{\mathrm{h}}\mathrm{C}_{\mathrm{h}}(\mathrm{T}_{1}-\mathrm{T}_{2}) = \mathrm{UA}\Delta t_{\mathrm{LMTD}}$$
(8-171)

Equating I and II in Equation 8-171

$$Mc \frac{dt}{d\theta} = W_{b}c(t'-t)$$
(8-172)

Rearranging Equation 8-172 gives:

$$t' = t + \frac{M}{W_{b}} \cdot \frac{dt}{d\theta}$$
(8-173)

The parameter S can be defined by

$$S = \frac{t'-t}{T_1-t} = \left(\frac{M}{W_b}\right) \left(\frac{1}{T_1-t}\right) \frac{dt}{d\theta}$$
(8-174)

The parameter R can be defined by equating II and III in Equation 171

$$R = \frac{T_{1} - T_{2}}{t' - t} = \frac{W_{b}c}{W_{h}C_{h}}$$
(8-175)

Rearranging Equation 8-174 gives:

$$\int_{t_{1}}^{t_{2}} \frac{dt}{T_{1} - t} = \frac{S.W_{b}}{M} \int_{0}^{\theta} d\theta$$
 (8-176)

Integrating from t_1 to t_2 , as the time passes from 0 to θ yields:

$$\ln\left(\frac{T_1 - t_1}{T_1 - t_2}\right) = \left(\frac{S.W_b}{M}\right)\theta$$
(8-177)

and

$$S = \frac{2(K_7 - 1)}{K_7 \{R + 1 + (R^2 + 1)^{0.5}\} - \{R + 1 - (R^2 + 1)^{0.5}\}}$$
(8-178)

where $K_7 = \exp\{(UA/W_b c)(R^2 + 1)^{0.5}\}$

A = heat transfer surface

- c = specific heat of batch liquid
- C_{h} = heating medium specific heat
- M = weight of batch liquid
- T_1 = initial temperature of heating medium
- t_1 = initial batch temperature
- $t_2 = final batch temperature$

U = overall heat transfer coefficient W_b = batch flow rate W_h = heating medium flow rate θ = time

Batch Heating and Cooling: External Heat Exchanger (2-4 Multipass Heat Exchangers)

The same procedure can be used as in 1-2 multipass heat exchangers. The equation for batch heating with external 2–4 multipass heat exchangers and non-isothermal heating media is

$$\ln\left(\frac{T_{1} - t_{1}}{T_{1} - t_{2}}\right) = \left(\frac{S.W_{b}}{M}\right)\theta$$
(8-177)

where

$$S = \frac{2(K_8 - 1)[1 + \{(1 - S)(1 - RS)\}^{0.5}]}{(K_8 - 1)(R + 1) + (K_8 + 1)(R^2 + 1)^{0.5}}$$
(8-179)

and

$$K_{8} = \exp\left\{\frac{UA}{2w_{b}c}(R^{2}+1)^{0.5}\right\}$$
(8-180)

Because S cannot be expressed explicitly, Equation 8-179 can be solved only by trial and error, assuming different values of S until an equality is attained. The equations for batch heating and cooling are the same as those developed for the 1-2 multipass exchangers, except that the value of S from Equation 8-179 replaces the value of S from Equation 8-178.

PROBLEMS AND SOLUTIONS

Problem 8-1

Kerosene leaves the bottom of a distillation column at 390°F and will be cooled to 200°F by crude oil from storage at 100°F and heated to 170°F. Calculate the number of shell passes required, the F-factor, and the corrected LMTD.

Hot kerosene inlet temperature $= 390^{\circ}F$

Hot kerosene outlet temperature $= 200^{\circ}F$

Crude oil inlet temperature	= 100°F
Crude oil outlet temperature	= 170°F

Solution

The computer program PROG81 calculates the required number of shell passes, the F-factor, the Log Mean Temperature Difference (LMTD), and the Corrected Mean Temperature Difference (CMTD). Table 8-21 shows the input data and computer output of the problem. The number of shells required is 1, the F-factor is 0.8917, the LMTD is 152.2°F and the corrected LMTD is 135.7°F.

Problem 8-2

Acetone (s = 0.79) at 250°F is to be sent to storage at 100°F and at a rate of 60,000 lb/hr. The heat will be received by 185,000 lb/hr of 100 percent acetic acid (s = 1.07) coming from storage at 90°F and heated to 150°F. Pressure drops of 10.0 psi are available for both fluids. Assuming that the fouling factor on the tube side is 0.001 and that on the shell side is 0.003, calculate the heat transfer coefficients for the tube and shell sides, the overall heat transfer coefficient for the exchanger, outside area of unit, and the heat transferred.

Available for the service are a large number of 1-2 exchangers having 21 1/4 in. ID shells with 270 tubes 3/4 in. OD, 14 BWG, 16 ft long

Table 8-21Input Data and Computer Output for the Corrected MeanTemperature Difference in a Shell and Tube Heat Exchanger

DATA81.DAT	
390.0 200.0 100.0 170.0 1.	
THE CORRECTED LMTD IN A SHELL AND TUBE	

HOT FLUID INLET TEMPERATURE, OF:	390.000
HOT FLUID INLET TEMPERATURE, oF:	390.000
HOT FLUID OUTLET TEMPERATURE, oF:	200.000
HOT FLUID INLET TEMPERATURE, OF:	390.000
HOT FLUID OUTLET TEMPERATURE, OF:	200.000
COLD FLUID INLET TEMPERATURE, OF:	100.000
HOT FLUID INLET TEMPERATURE, oF:	390.000
HOT FLUID OUTLET TEMPERATURE, oF:	200.000
HOT FLUID INLET TEMPERATURE, OF:	390.000
HOT FLUID OUTLET TEMPERATURE, OF:	200.000
COLD FLUID INLET TEMPERATURE, OF:	100.000
HOT FLUID INLET TEMPERATURE, OF:	390.000
HOT FLUID OUTLET TEMPERATURE, OF:	200.000
COLD FLUID INLET TEMPERATURE, OF:	100.000
COLD FLUID OUTLET TEMPERATURE, OF:	170.000
HOT FLUID INLET TEMPERATURE, OF:	390.000
HOT FLUID OUTLET TEMPERATURE, OF:	200.000
COLD FLUID INLET TEMPERATURE, OF:	100.000
COLD FLUID OUTLET TEMPERATURE, OF:	170.000
NUMBER OF SERIES EXCHANGER SHELLS:	1.
HOT FLUID INLET TEMPERATURE, OF:	390.000
HOT FLUID OUTLET TEMPERATURE, OF:	200.000
COLD FLUID INLET TEMPERATURE, OF:	100.000
COLD FLUID OUTLET TEMPERATURE, OF:	170.000
NUMBER OF SERIES EXCHANGER SHELLS:	1.
THE NUMBER OF SHELLS REQUIRED:	1.

and laid on 1-in. square pitch. The bundles are arranged for two tube passes with segmental baffles spaced 5 in. apart.

Physical property data:

Physical property	Acetone	Acetic Acid
Viscosity, cP	0.20	0.85
Density, lb/ft ³	49.296	66.768
Specific heat, Btu/lb.°F	0.63	0.51
Thermal conductivity, Btu/lb.ft ² (°F/ft)	0.095	0.098

Solution

Exchanger:

Hot Fluid (Acetone)	Cold Fluid (Acetic Acid)
Shell side	Tube side
ID=21 1/4 in.	Number and length = $270, 16'0''$
Baffle space $= 5in$.	OD, BWG, pitch = 3/4in., 14 BWG, 1 in. square.
Passes = 1	Passes = 2

Heat Balance:

Heat lost by acetone to storage = heat gained by acetic acid from storage

that is,

 $W.C_{p}(T_{2} - T_{1}) = w.c_{p}(t_{2} - t_{1})$ (60,000)(0.57)(250 - 100) = w(0.51)(150 - 90) $w \approx 168,000 \text{ lb/hr}$

The computer program PROG82 rates the shell and tube heat exchanger using 168,000 lb/hr of acetic acid in the tube side. Table 8-22 gives the input data and computer output of the heat exchanger design. A minimum of three shell passes is required to permit the transfer of heat with the temperatures given by the process. The overall heat transfer coefficient is 79 Btu/hrft²°F, and the heat transfer surface is 2545 ft². Three exchangers are sufficient for the heat transfer, although the pressure

Table 8-22 Input Data and Computer Output for the Rating of a Shell and Tube Heat Exchanger

DATA82.DAT						
ACETIC 270.0	ACID 0.584	0.75	1.0			
S 16.0 150.0 0.098	2.0 0.51 168000.	0.001 0.85	90.0 66.768			
ACETONE 1.0 0.003 0.20	21.25 250.0 49.296	38.0 100.0 0.095	5.0 0.63 60000.0			

SHELL AND TUBE HEAT EXCHANGER RATING

	TUBE SIDE	
FLUID TYPE: FLUID TYPE: FLUID FLOW RATE , lb/hr.: SPECIFIC HEAT CAPACITY, Btu/lb.oF: FLUID DENSITY, lb/ft^3: FLUID VISCOSITY, cP: FLUID VISCOSITY, cP: FLUID INLET TEMPERATURE, oF: FLUID OUTLET TEMPERATURE, oF: INSIDE DIAMETER, inch: OUTSIDE DIAMETER, inch: TUBE PITCH, inch: TUBE PITCH, inch: TUBE LENGTH, ft.: NUMBER OF TUBES: NUMBER OF PASSES: NUMBER OF BAFFLES: BAFFLE SBACIMC inch:	ACETIC ACID	ACETONE
FLUID FLOW RATE , lb/hr.:	168000.	60000.
SPECIFIC HEAT CAPACITY, Btu/lb.oF:	.510	.630
FLUID DENSITY, 1b/ft^3:	66.768	49.296
FLUID VISCOSITY, CP:	.850	.200
FLUID THERMAL COND. Btu/ft.h.oF:	.098	.095
FLUID INLET TEMPERATURE, OF:	90.	250.
FLUID OUTLET TEMPERATURE, oF:	150.	100.
INSIDE DIAMETER, inch:	.584	21.250
DUTSIDE DIAMETER, inch:	.750	
TUBE PITCH, inch:	1.000	SQUARE PITCH
TUBE LENGTH, ft.:	16.000	
IUMBER OF TUBES:	270.	
UMBER OF PASSES:	6.	з.
NUMBER OF BAFFLES: SAFFLE SPACING, inch: PLUID VELOCITY, ft/sec.: REYNOLDS NUMBER: PRICTION FACTOR (ft^2/in^2):		38.
BAFFLE SPACING, inch:		5.0
FLUID VELOCITY, ft/sec.:	2,783	1.833
REYNOLDS NUMBER:	15828.	53072.
PRANDTL NUMBER:	10.7048	3.2097
FRICTION FACTOR (ft^2/in^2):	.00024	.00153
<pre>{EAT COEFF., Btu/hr.ft^2.oF:</pre>	211.8	253.4
PRESSURE DROP, psi:	1.5682	3.4464
FOULING FACTOR:	.0010	.0030
APPROACH FACTOR, F:	.8749	
RANDTL NUMBER: FRICTION FACTOR (ft^2/in^2): HEAT COEFF., Btu/hr.ft^2.oF: RESSURE DROP, psi: OULING FACTOR: PPROACH FACTOR, F: OG MEAN TEMP. DIFF. oF:	39.09	
CORRECTED MEAN TEMP. DIFF. oF:	34.20	
OVERALL HEAT COEFF., Btu/hr.ft^2.oF:	79.0	
OUTSIDE AREA OF UNIT, ft^2:	2544.7	
DUTSIDE AREA OF UNIT, ft^2: HEAT LOAD ON UNIT, Btu/hr:	5140800.	
HEAT TRANSFERRED, Btu/hr:	6870014.	

drop on the shell side is higher than the permissible pressure drop. Fewer exchangers will be inadequate for the service.

Problem 8-3

Petroleum distillate oil (28° API) is to be cooled in a double pipe finned tube exchanger with cooling-tower water under the following conditions:

Tube side: Water flow = 23,310 lb/hr $T_{in} = 80^{\circ}F$ $T_{out} = 120^{\circ}F$ Fol = 0.003 $\mu = 0.72 \text{ cP}$ $K = 0.366 \text{ Btu/fth}^{\circ}F$ $C_p = 1 \text{ Btu/lb}^{\circ}F$ Density = 62.3 lb/ft³ Number of passes = 5

Shell side: Oil flow = 18,000 lb/hr $T_{in} = 250^{\circ}F$ $T_{out} = 150^{\circ}F$ Fol = 0.001 $\mu = 2.45cP$ K = 0.074 Btu/ft hr°F $C_p = 0.518$ Btu/lb°F Density = 45 lb/ft³ Number of passes = 1

Exchanger size: Shell side: 3 in. (3.068 in. ID, 3.5 in. OD) Tube side: 1 1/2 in. (1.61 in. ID, 1.9 in. OD) Tube length: 12 ft Fins: 24, 0.5 in. high x 0.035 in. wide Thermal conductivity of tube material

 $K = 25 \text{ Btu/fthr}^{\circ}\text{F}$

Pressure drops of 10 psi are allowable on both streams, and the fouling factors of 0.001 for the distillate oil and 0.003 for the water are required. Calculate the overall heat transfer coefficient, and the exchanger fin efficiency.

Solution

Passes = 5
Tube length = $12'0''$
ID = 1.61 in. $OD = 1.9$ in.
Tube side
Cold Fluid (Water)

Physical property data:

Physical property	Oil	Water
Viscosity, cP	2.45	0.72
Density, lb/ft ³	45.0	62.3
Specific heat, Btu/lb.°F	0.518	1.00
Thermal conductivity, Btu/lbft2(°F/ft)	0.074	0.366

Heat Balance:

Heat lost by oil = heat gained by water

that is,

 $Q = W.Cp.(T_1 - T_2) = w.cp.(t_2 - t_1)$ Q = (18000)(0.518)(250 - 150) = w(1)(120 - 80)932400 = 40ww = 23310 lb/hr

The flow of water required to cool the oil is 23310 lb/hr. The computer program PROG83 rates the double pipe heat exchanger using either fin tubes or bare tubes. Table 8-23 lists the input data and output of the computer program. The fin efficiency is 43.3 percent and the computed overall heat transfer coefficient is 27.3 Btu/hrft²°F. The pressure drops have not been exceeded, and the double pipe exchanger will be satisfactory for the service.

Problem 8-4

Calculate a preliminary air-cooler design using the following data.

A light hydrocarbon liquid is to be cooled from 170° F to 135° F in an air cooler. The heat transfer rate is 5.0 MM Btu/hr, and the estimated overall heat transfer is 60 Btu/hr ft²°F. The design dry bulb air temperature is 85°F, and the tube length is 40ft.

Solution

The computer program PROG84 calculates the air cooled heat exchanger area, the face area and the fan horsepower. Table 8-24 shows both the input data and computer output of the design calculations. The exchanger area is 1602 ft^2 and the LMTD is 52°F . Face area of the bundle

Table 8-23
Input Data and Computer Output for the Double Pipe
or Longitudinal Finned Tube Heat Exchanger

	рат.			
DATA83	• DAT			
WATER				
LIQUID				
1.61	1.9	23310.0		
1.0	120 0	0.003		
60.0	120.0	23310.0 0.366 0.003 5		
OIL	12.0	5		
FINS				
LIQUID				
24				
3.068	3.5	18000.0		
0.518	2.45	0.074		
250.0	150.0	0.001		
0.5	45 0	18000.0 0.074 0.001		
25.0	43.0	1		
		DOUBLE PIPE HEAT EXCHANGER	RATING	
		USING BARE-TUBES/LONGITUDI		
		FORCED CONVECTION WITH NO	CHANGE OF PHA	SE
*****	******			
		E FLOW: ATE, 1b/hr.: T CAPACITY, Btu/Ib.oF: Y, 1b/ft^3.: ITY, cP: L COND., Btu/Ib.oF: TEMPERATURE, oF: TEMPERATURE, oF: TER, inch: ETER, inch: OR: ft.: BE PASSES:		SHELL SIDE
FLUID	NAME:		WATER	OIL
TYPE (OF PHAS	E FLOW:	LIQUID	LIQUID
FLUID	FLOW R	ATE, lb/hr.:	23310.	18000.
SPECI	FIC HEA	r CAPACITY, Btu/15.of:	1.000	.518
FLUID	DENSIT	Y, ID/IC^3.:	62.300	2 450
FLUID	THERMA	L COND. Btu/Tb.oF	. 366	.074
FLUID	INLET	TEMPERATURE. OF:	80.000	250.000
FLUID	OUTLET	TEMPERATURE, oF:	120.000	150.000
INSID	E DIAME	TER, inch:	1.610	3.068
OUTSI	DE DIAM	ETER, inch:	1.900	3.500
FOULI	NG FACT	OR:	.003	.001
TUBE	LENGTH,	ft.:	12.000	
NUMBE	R OF TU	BE PASSES:	24	
	EIGHT,		.500	
		S, inch:	.035	
		. TUBE MATERIAL, Btu/hr.ft.oF:		
		TY, ft/sec.:	7.351	3.867
	LDS NUM	BER:	126958.	3759.
	TL NUMB	ER: TOR (ft^2/in^2):	4.7607 .000031	41.503 .000042
		P, psi.:	2.9424	.5943
	ODULUS:	P, psi.:	2.258	: 3943
	FFICIEN	CV &•	43.335	
		RFACE EFFICIENCY, %:	44.486	
TOC N	EAM DEM	D DIEE OF.	96,92	
HEAT	TRANSFE	R FILM COEFF. CORRECTED		
FOR F	OULING:	, Btu/hr.ft^2.oF:	264.33	107.04
OVERA	LL HEAT	, Btu/hr.ft^2.oF: TRANSFER COEFF., Btu/hr.ft^2.	27.26	
HEAT	TOWD ON	UNIT, DCU/III	932400.0 352.932	
HEAT	TRANSFE	R SURFACE ft^2.:	352.932	

is 283ft², and the fan horsepower is 25.2 bhp. The required air flow is 164266 std ft³/min.

Problem 8-5

Determine the number of tracers required to maintain 100 ft of 8-in. (Schedule 40) process line (ID = 7.981 in.) at 500°F. Hot tracing

Table 8-24 Input Data and Computer Output for An Air-Cooled Heat Exchanger Design

DATA84.	DAT
---------	-----

170.0 60.0	135.0 5000000.0	85.0 40.0	
		AN AIR-COOLED HEAT EXCHANGER DESIGN	
*****	********	**************************************	****
			170.0
		S FLUID TEMPERATURE, oF:	135.0
		PERATURE, OF:	85.0
		TRANSFER COEFFICEINT, Btu/hr.ft^2.oF:	
HI	EAT LOAD, Bt	u/hr.:	5000000.
	JBE LENGTH,		40.0
NU	MBER OF TUB	E ROWS:	4.5
RA	TIO OF BARE	-TUBE SURFACE TO FACE AREA OF BUNDLE:	57
FA	CE VELOCITY	OF AIR, ft./min.:	580.740
		OUTLET TEMPERATURE, oF:	105.250
		MEAN TEMPERATURE DIFFERENCE, oF:	57.058
BA	RE-TUBE SUR	FACE AREA BASED ON TUBE OD, ft^2:	1460.5
FA	CE AREA OF	BUNDLE, ft^2:	257.9
CA	LCULATED OU	TLET AIR TEMPERATURE, OF:	115.916
Tł	E DIFFERENC	E BETWEEN THE CALCULATED AIR	
		ATURE AND THE ESTIMATED AIR OUTLET	
		S GREATER THAN 0.5	TAO-TAO1 >0.5
EF	FECTIVE LOG	MEAN TEMPERATURE DIFFERENCE, oF:	52.015
BA	RE-TUBE SUR	FACE AREA BASED ON TUBE OD, ft^2:	1602.1
r P	CE AREA UF .	BUNDLE, IT^2:	282.9
		ILET AIR TEMPERATURE, OF:	113.184
CA	LCULATED BA	RE-TUBE SURFACE AREA	
BA	SED ON TUBE	OUTSIDE DIAMETER, ft^2:	1602.1
FA	CE AREA OF	BUNDLE, ft^2:	282.9
		TUBES, std.ft ³ /min.:	164266.
	N HORSEPOWER		25.2
	R-COOLER WE		22169.8
TU	BE BUNDLE W	IDTH, ft.:	7.1

medium is available at 630°F and has a heat capacity of 0.53 Btu/lb°F. The process line is covered with 2.5 in. of insulation, and this insulation has a thermal conductivity of 0.033 Btu/(hr)(ft²)(°F/ft). Design for 0°F air temperature and 20 mph winds. Use 1/2 in. tracers, the hot medium outlet temperature is 550°F. Allow approximately 1 1/4 in. between pipe and insulation to accommodate a 1/2 in. in tracer line and heat transfer cement. Assuming the trial film heat transfer coefficient is 4.0 Btu/hrft² °F, and the thermal conductances tracer to 1/2 in. pipe are a = 0.393 and b = 4.58 Btu/hr °F(ft of pipe).

Solution

The computer program PROG85 calculates the heat tracer requirements and heat loss for an insulated pipe line either with tracing or without tracing. Table 8-25 lists the input data and computer output of the

Table 8-25 Input Data and Computer Output for Tracer Requirements for Pipelines

DATA85	חאת
DATAGO	+ DAT

100.0	500.0	630.0	
550.0	0.0	7.981	
1.25	2.5	0.033	
0.53	4.0	0.393	4.58
0.53	4.0	0.393	4.58

HEAT TRACER REQUIREMENTS FOR PIPELINES AND	
HEAT LOSS FROM AN INSULATED PIPELINE WITH TRACIN	IC.

TOTAL PIPELINE LENGTH, ft.:	100.0
TEMPERATURE IN PIPE, oF:	500.0
	630.0
HOT MEDIUM OUTLET TEMPERATURE, OF:	550.0
	.0
OUTSIDE DIAMETER OF PIPE, in.:	7.981
	1.25
INSULATION THICKNESS, in.:	2.50
THERMAL CONDUCTIVITY OF INSULATION, Btu/hr.ft^2(oF/1	ft .033
SPECIFIC HEAT OF HOT MEDIUM, Btu/Ib.oF:	.530
FILM HEAT TRANSFER COEFFICIENT TO AIR	
CORRECTED FOR WIND, Btu/ft^2.hr.oF:	4.000
THERMAL CONDUCTANCE TRACER TO PIPE	
	.393
THERMAL CONDUCTANCE TRACER TO PIPE	
WITH CEMENT, Btu/hr.oF.ft of pipe: OUTSIDE SURFACE TEMPERATURE OF INSULATION, oF: THE COMBINED CONVECTIVE AND RADIATIVE	4.580
OUTSIDE SURFACE TEMPERATURE OF INSULATION, OF:	16.972
THE COMBINED CONVECTIVE AND RADIATIVE	
HEAT TRANSFER COEFFICIENT, Btu/h.oF.ft^2:	1.330
WIND FACTOR:	2.807
HEAT TRANSFER COEFFICIENT TO AIR	
CORRECTED FOR THE WIND, Btu/hr.ft^2.oF:	3.734
HEAT LOSS PER FOOT OF PIPE, Btu/hr.ft:	252.934
CORRECTED FOR THE WIND, Btu/hr.ft^2.oF: HEAT LOSS PER FOOT OF PIPE, Btu/hr.ft: TOTAL HEAT LOSS FROM PIPELINE, Btu/hr.: FLOW RATE OF HOT MEDIUM, 1b/hr.:	25293.430
FLOW RATE OF HOT MEDIUM, 1b/hr.:	596.543
THE NUMBER OF TRACERS WITHOUT HEAT TRANSFER CEMENT:	7.15
THE NUMBER OF TRACERS WITHOUT HEAT TRANSFER CEMENT: THE NUMBER OF TRACERS WITH HEAT TRANSFER CEMENT:	.61
UPLE LOGG PROV IN THEME PEOP STOP STOP STOP	
HEAT LOSS FROM AN INSULATED PIPE WITHOUT TRACING	
OUTSIDE SURFACE TEMPERATURE OF INSULATION, OF:	15.202
THE COMBINED CONVECTIVE AND RADIATIVE	

THE COMBINED CONVECTIVE AND RADIATIVE	
HEAT TRANSFER COEFFICIENT, Btu/h.oF.ft^2:	1.344
WIND FACTOR:	2.808
HEAT TRANSFER COEFFICIENT TO AIR	
CORRECTED FOR THE WIND, Btu/hr.ft^2.oF:	3.774
HEAT LOSS PER FOOT OF PIPE, Btu/hr.ft:	206.652
TOTAL HEAT LOSS FROM PIPELINE, Btu/hr.:	20665.240

problem. The approximate number of tracers without the heat transfer cement of an 8-inch (Schedule 40) process line is 7. The heat transfer cement reduces the number required to 1. The circulation rate of the tracing medium is 596.5 lb/hr, and the heat lost from 100 ft. of pipe line is 25293.4 Btu/hr.

Problem 8-6 (Internal Coil, Isothermal Heating Medium)

A tank containing 55,000 lb (24947.6 kg) of material with a specific heat of 0.5 Btu/lb °F (2.093 kJ/kg.K) is heated from 72°F (295.4K) to 260°F (399.8K). The tank contains a heating coil with a heat transfer surface of 110 ft² (10.22m²), and the overall heat transfer coefficient from the coil to the tank contents is 150 Btu/(hr)(ft²)(°F)(851.7 W/m².K). Calculate the time required to heat the tank contents with steam condensing at 350°F (449.8K).

Solution

When heating a batch reactor with an internal coil under isothermal condition, Equation 8-135 applies:

$$\ln\left(\frac{T_1 - t_1}{T_1 - t_2}\right) = \frac{UA}{Mc} \cdot \theta$$
(8-135)

The time, θ , required to heat the tank contents with steam condensing at 350°F is,

$$\theta = \frac{\ln\left(\frac{T_1 - t_1}{T_1 - t_2}\right)}{\left(\frac{UA}{Mc}\right)}, \text{ hr}$$
$$= \frac{\ln\left(\frac{350 - 72}{350 - 260}\right)}{\left(\frac{150 \times 110}{55,000 \times 0.5}\right)}$$
$$= 1.87 \text{ hr}$$

Problem 8-7 Batch Cooling: External Heat Exchanger (Counterflow), Non-isothermal Cooling Medium

For the tank described in PROBLEM 8-6, calculate the time required to cool the batch from $260^{\circ}F(399.8K)$ to $150^{\circ}F(338.7K)$, if an external

heat exchanger with a heat transfer surface of 250ft² (23.23m²) is available. The batch material is circulated through the exchanger at the rate of 30,000 lb/hr (13608 kg/h). Cooling water is available at a temperature of 85°F (302.6K) and a flow rate of 10,000 lb/hr (4536 kg/h). The overall heat transfer coefficient in the heat exchanger is 250 Btu/hr ft² °F (1419.5 W/m² K).

Solution

Equation 8-169 applies when cooling a batch with an external heat exchanger and a non-isothermal cooling medium.

$$\ln\left(\frac{\mathbf{T}_{1}-\mathbf{t}_{1}}{\mathbf{T}_{1}-\mathbf{t}_{2}}\right) = \left(\frac{\mathbf{K}_{5}-1}{\mathbf{M}}\right) \left(\frac{\mathbf{W}_{b}\mathbf{W}_{c}\mathbf{C}_{c}}{\mathbf{K}_{5}\mathbf{W}_{c}\mathbf{C}_{c}-\mathbf{W}_{b}\mathbf{c}}\right) \boldsymbol{\theta}$$
(8-169)

where

$$K_{5} = \exp\left\{UA\left(\frac{1}{W_{b}c} - \frac{1}{W_{c}C_{c}}\right)\right\}$$

Using the values in Table 8-26, K₅ is calculated as follows:

$$K_{5} = \exp\left\{250 \times 250 \left(\frac{1}{30,000 \times 0.5} - \frac{1}{10,000 \times 1.0}\right)\right\}$$
$$= 0.1245$$

The time, θ , required to cool the batch is determined from Equation 8-169,

$$\ln\left\{\frac{260-85}{260-85}\right\} = \left(\frac{0.125-1}{55,000}\right)$$
$$\times \left(\frac{30,000(10,000)(1.0)}{0.1245(10,000)(1.0)-(30,000)(0.5)}\right)\theta$$

 $\theta = 2.854$ hr

Table 8-26

A =250	$W_{b} = 30,000$	M =55,000	C =0.5	$T_2 = 150^{\circ}F$
U =250	$W_{c} = 10,000$	$C_{c} = 1.0$	$T_1 = 260^{\circ}F$	$t_1 = 85^{\circ}F$

Problem 8-8

8000 gal (30.28m³) of liquid hydrocarbon under pressure at 350°F (449.8K) is required for a batch extraction process. The storage temperature of the liquid is 100°F (310.9K). Available as a heating medium is a 12,000 lb/hr (5443.2 kg/h) 28° API oil stream at a temperature of 400°F (477.6K). A pump connected to the batch is capable of circulating 50,000 lb/hr (22680.0 kg/h) of the hydrocarbon. Available for the service is 400 ft² (37.16m²) of clean double pipe exchanger surface that gives a U_e of 50 Btu/hrft²°F (283.9 W/m².K) in counterflow streams.

Calculate for the flow rates:

- 1. The time required to heat the agitated batch using the double pipe countercurrent flow exchanger.
- 2. The time required using a 1-2 exchanger with the same surface and coefficient.
- 3. The time required using a 2-4 exchanger with the same surface and coefficient.

Physical property data:

Liquid hydrocarbon Specific gravity: 0.88 Specific heat capacity: 0.48 Btu/lb °F (2.01kJ/kg.K)

28° API oil: Specific heat capacity: 0.6 Btu/lb °F (2.51 kJ/kg.K)

Solution

Volume of liquid hydrocarbon in the batch = 8000 gal Specific gravity = 0.88

Mass (M) of liquid hydrocarbon in the batch = $8000 \times 0.88 \times 8.3$	4
= 58713.6	
≈ 59,000 lb.	

V = 8000	$W_{h} = 12,000$	$C_{h} = 0.6$	A = 400	$t_1 = 100^{\circ}F$
$W_b = 50,000$	$U_{c} = 50$	c = 0.48	$T_1 = 400^{\circ}F$	$t_2 = 350^{\circ}F$

Table 8-27

(1) The time required to heat the agitated batch using the double pipe countercurrent exchanger can be determined from Equation 8-170.

$$\ln\left(\frac{\mathbf{T}_{1}-\mathbf{t}_{1}}{\mathbf{T}_{1}-\mathbf{t}_{2}}\right) = \left(\frac{\mathbf{K}_{6}-1}{\mathbf{M}}\right)\left(\frac{\mathbf{W}_{b}\mathbf{W}_{h}\mathbf{C}_{h}}{\mathbf{K}_{6}\mathbf{W}_{h}\mathbf{C}_{h}-\mathbf{W}_{b}\mathbf{c}}\right)\boldsymbol{\theta}$$

where

$$\begin{split} \mathbf{K}_{6} &= \exp\left\{ \mathrm{UA}\left(\frac{1}{\mathrm{W}_{b}\mathrm{c}} - \frac{1}{\mathrm{W}_{h}\mathrm{C}_{h}}\right) \right\} \\ \mathbf{W}_{b}\mathrm{c} &= (50,000)(0.48) \\ &= 24,000 \\ \mathbf{W}_{h}\mathrm{C}_{h} &= (12,000)(0.60) \\ &= 7,200 \\ \mathbf{K}_{6} &= \exp\left\{ (50 \times 400) \left(\frac{1}{24,000} - \frac{1}{7,200}\right) \right\} \\ &= 0.143 \\ \ln\left(\frac{400 - 100}{400 - 300}\right) &= \left(\frac{0.143 - 1}{59,000}\right) \left(\frac{(50,000)(12,000)}{(0.143)(7,200) - 24,000}\right) \theta \\ \mathrm{h} &= 7.87 \,\mathrm{hr} \end{split}$$

(2) Equation 8-177 applies for the time to heat the agitated batch using a 1-2 exchanger.

$$\ln\left(\frac{T_1 - t_1}{T_1 - t_2}\right) = \left(\frac{S.W_b}{M}\right)\theta$$
(8-177)

and

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$$S = \frac{2(K_7 - 1)}{K_7 \{R + 1 + (R^2 + 1)^{0.5}\} - \{R + 1 - (R^2 + 1)^{0.5}\}}$$
(8-178)

٠

where

$$\mathbf{K}_{7} = \exp\left\{\frac{\mathbf{U}\mathbf{A}}{\mathbf{W}_{b}\mathbf{c}}\left(\mathbf{R}^{2}+1\right)^{0.5}\right\}$$

and

$$R = \frac{W_{b}c}{W_{h}C_{h}}$$

$$= \frac{24,000}{7,200}$$

$$= 3.33$$

$$K_{7} = \exp\left\{\frac{(50)(400)}{24,000} (3.33^{2} + 1)^{0.5}\right\}$$

$$= 18.127$$

Substituting the value of K_7 into Equation 8-178 gives,

$$S = \frac{2(18.127 - 1)}{18.127 \{ 3.33 + 1 + (3.33^2 + 1)^{0.5} \} - \{ 3.33 + 1 - (3.33^2 + 1)^{0.5} \}}$$

= 0.2435

The value of S in Equation 8-177 gives

$$\ln\left(\frac{400-100}{400-350}\right) = \left(\frac{0.2435 \times 50,000}{59,000}\right)\theta$$

 $\theta = 8.68 \text{ hr}$

(3) Equations 8-177, 8-179, and 8-180 apply for the time to heat the agitated batch using a 2-4 exchanger.

$$S = \frac{2(K_8 - 1)\left\{1 + \left[(1 - S)(1 - RS)\right]^{0.5}\right\}}{(K_8 - 1)(R + 1) + (K_8 + 1)(R^2 + 1)^{0.5}}$$
(8-179)

and

$$K_{8} = \exp\left\{\frac{UA}{2W_{b}c}(R^{2}+1)^{0.5}\right\}$$
(8-180)

Equation 8-179 is an implicit equation involving S on both sides, and can be solved by trial and error until an equality is reached. Rearranging Equation 8-179 gives

LHS RHS

$$S\{(K_{8} - 1)(R + 1) + (K_{8} + 1)(R^{2} + 1)^{0.5}\}$$

$$= 2(K_{8} - 1)\{1 + [(1 - S)(1 - RS)]^{0.5}\}$$
(8-181)

$$K_{8} = \exp\{\left(\frac{50 \times 400}{2 \times 50,000 \times 0.48}\right)(3.33^{2} + 1)^{0.5}\}$$

$$= 4.2576$$

$$R = 3.33$$

Substituting S = 0.262 in Equation 8-177 gives

$$\ln\left(\frac{400-100}{400-350}\right) = \left(\frac{0.262\times50,000}{59,000}\right)\theta$$

$$\theta = 8.07 \, \mathrm{hr}$$

Table 8-28

S	LHR	RHS
0.260	8.418	8.566
0.261	8.45	8.539
0.262	8.588	8.573
0.263	8.515	8.485

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PROGRAM PROG81

c c	THIS PROGRAM CALCULATES THE CORRECTED L.M.T.D. IN SHELL-AND-TUBE HEAT EXCHANGERS
00000000000	 THE PROGRAM CALCULATES: 1. THE IDEAL AND TRUE MEAN TEMPERATURE DIFFERENCE FOR ONE OR MORE EXCHANGER IN SERIES. 2. THE PROGRAM WILL INCREASE THE NUMBER OF EXCHANGER SHELL UNTIL A 'REASONABLE' F-FACTOR IS OBTAINED THUS ELIMINATING THE NEED FOR THE CHARTS AND GRAPHS NORMALLY USED TO CALCULATE F-FACTORS FOR SHELL-AND-TUBE HEAT EXCHANGERS. THE PROGRAM IS BASED ON THE FOLLOWING ASSUMPTIONS:
	 THE HOT AND COLD FLUIDS FLOW COUNTERCURRENT TO EACH OTHER. THE OVERALL HEAT TRANSFER COEFFICIENT (U) IS CONSTANT THROUGHOUT THE HEAT EXCHANGER. THE FLOW RATE OF EACH FLUID IS CONSTANT. THE SPECIFIC HEAT OF EACH FLUID IS CONSTANT. THERE IS NO CONDENSATION OF VAPOR OR BOILING OF LIQUID IN ANY PART OF THE EXCHANGER. HEAT LOSSES ARE NEGLIGIBLE. THERE IS EQUAL HEAT TRANSFER SURFACE IN EACH PASS. THE TEMPERATURE OF THE SHELL-SIDE FLUID IN ANY SHELL-SIDE PASS IS UNIFORM OVER ANY CROSS-SECTION.
000000000000000000000000000000000000000	<pre>thit = HOT FLUID INLET TEMPERATURE, oF. TH1 = HOT FLUID OUTLET TEMPERATURE, oF. TC1 = COLD FLUID INLET TEMPERATURE, oF. TC2 = COLD FLUID OUTLET TEMPERATURE, oF. N = NUMBER OF SERIES EXCHANGER SHELLS REQUIRED. LMTD = LOG MEAN TEMPERATURE DIFFERENCE. F = CORRECTION FACTOR. DELT1 = TH2-TC1, oF. DELT2 = TH1-TC2, oF. DELT = LMTD. ************************************</pre>
	REAL LMTD, N OPEN (UNIT=3, FILE='DATA81.DAT', STATUS='OLD', ERR=18) OPEN (UNIT=1, FILE='PRN')
C C C C C C	READ THE HOT FLUID INLET TEMPERATURE, OF: TH1 READ THE HOT FLUID OUTLET TEMPERATURE, OF: TH2 READ THE COLD FLUID INLET TEMPERATURE, OF: TC1 READ THE COLD FLUID OUTLET TEMPERATURE, OF: TC2 READ THE INITIAL NUMBER OF SERIES EXCHANGER SHELLS REQUIRED: N READ (3, *, ERR=19) TH1, TH2 READ (3, *, ERR=19) TC1, TC2 READ (3, *, ERR=19) N GO TO 10

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18 100	WRITE (*, 100) FORMAT (6X, 'DATA FILE DOES NOT EXIST') GO TO 999
19 110	WRITE (*, 110) Format (6X, 'Error message in the data value') Go to 999
10 120	WRITE (1, 120) FORMAT (///,10X,'THE CORRECTED LMTD IN A SHELL AND TUBE HEAT',\)
130	WRITE (1, 130) FORMAT (' EXCHANGER',/1H ,78(1H*))
140 *	WRITE (1, 140) TH1, TH2 FORMAT(6X,'HOT FLUID INLET TEMPERATURE, oF:', T60, F9.3,/,6X, 'HOT FLUID OUTLET TEMPERATURE, oF:',T60, F9.3)
150 *	<pre>WRITE (1, 150) TC1, TC2 FORMAT(6X,'COLD FLUID INLET TEMPERATURE, oF:',T60, F9.3,/,6X,</pre>
160	WRITE (1, 160) N FORMAT(6X,'NUMBER OF SERIES EXCHANGER SHELLS:',T60,F4.0)
	P = (TC2-TC1)/(TH1-TC1) R = (TH1-TH2)/(TC2-TC1) RSQRT = (R**2+1.)**0.5
15	<pre>IF (R .EQ. 1.0) THEN GO TO 25 ELSE VAL1 = 1.0-((P*R-1.0)/(P-1.0))**(1./N) VAL2 = R-((P*R-1.0)/(P-1.0))**(1./N) P1 = VAL1/VAL2 ENDIF</pre>
	VAL3 = (2.0/P1)-1.0-R+RSQRT VAL4 = (2.0/P1)-1.0-R-RSQRT
	A = VAL3/VAL4
	IF (A .LT. 0.0) THEN GO TO 30 ELSE VAL5 = (RSQRT/(R-1.0))*ALOG((1.0-P1)/(1.0-P1*R)) F = VAL5/ALOG(A) ENDIF
	GO TO 40
30	N = N+1
	GO TO 15
25	P2 = P/(N-P*(N-1.)) VAL6 = (2.0/P2)-2.0+RSQRT

	VAL7 = (2.0/P2)-2.0-RSQRT B = VAL6/VAL7 IF (B .LT. 0.0) THEN GO TO 30 ELSE VAL8 = (RSQRT*P2)/(1.0-P2) F = VAL8/ALOG(B) ENDIF
40	IF (F .GT. 0.75) THEN GO TO 45 ELSE N = N+1 GO TO 15 ENDIF
45	DELT1 = TH1-TC2 DELT2 = TH2-TC1 IF (DELT1 .EQ. DELT2) THEN GO TO 50 ELSE
с	CALCULATE THE LOG MEAN TEMPERATURE DIFFERENCE
	LMTD = (DELT1-DELT2)/ALOG(DELT1/DELT2) ENDIF GO TO 55
50	DELT = LMTD
с	CALCULATE THE CORRECTED LOG MEAN TEMPERATURE DIFFERENCE
55	$CLMTD = F \star LMTD$
170 * *	<pre>WRITE (1, 170) N, F, LMTD, CLMTD FORMAT (6X,'THE NUMBER OF SHELLS REQUIRED:',T60, F4.0,/,6X, 'THE F- FACTOR:',T60, F8.4,/,6X, 'THE LOG MEAN TEMPERATURE DIFFERENCE,oF:',T60,F8.4,/,6X, 'THE CORRECTED LMTD, oF:',T60, F8.4)</pre>
180	WRITE (1, 180) FORMAT (1H ,78(1H-))
с	FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR(12)
999	CLOSE (3,STATUS='KEEP') CLOSE (1) STOP END

PROGRAM PROG82

~		****	******	*******	*****
á		OGRAM RATES A SHELL AND '			
0		A SINGLE SHELL PASS AND A			
3				TWO TUBE	
C		OR TWO OR MORE SHELL PAS			
Ċ.	THE PRO	GRAM ASSUMES THAT THERE	IS NO CHANGE OF	PHASE.	
С					
С		GRAM USES DEVELOPED CORR			
С	IN BOTH	THE SHELL AND TUBE SIDE	S. HEDRICK'S CO	RRELATIO	N
с	(Chem.	Eng. June, 1990) FOR EST	IMATING THE HEAT	TRANSFE	R
c	COEFFIC	IENT FOR TUBES IN THE TR	ANSITION REGION	IS EMPLO	YED
c		PROGRAM.			
c	******	******	*****	******	* * * * *
	NOVENCI		TUBE SIDE		
с	NOMENCL	AIURE:	IUBE SIDE		
С					FTS
с	1.	FLUID TYPE			
С	2.	NUMBER OF TUBES		=	NT
с	3.	TUBE INSIDE DIA., inch		-	TID
С	4.	TUBE OUTSIDE DIA., inch		=	TOD
С	5.	TUBE PITCH, inch		=	TP
С	6.	PITCH TYPE (TRIANGLE/SQ	UARE)	=	T/S
С	7.	TUBE LENGTH, ft.		=	TL
С	8.	NUMBER OF PASSES		=	TNP
с	9.	FOULING FACTOR		#	TFF
с	9. 10. 11.	INLET TEMPERATURE, OF		=	TT1
С	11.	OUTLET TEMPERATURE, oF		*	TT2
с	12.	SPECIFIC HEAT CAPACITY	(Btu/lboF)	=	TSHC
С	13.	VISCOSITY, cP		=	TVIS
С	14.	DENSITY, 1b/ft^3.		=	TDEN
С	15.	DENSITY, 15/ft^3. FLUID THERMAL CONDUCTIV	ITY, (Btu/ft.h.o	F)=	TFK
С	16.			=	T₩
С		. ,			
c		NOMENCLATURE:	SHELL SIDE		
č					
c	1.	FLUID TYPE		=	FSS
č	2.	NUMBER OF SHELL PASSES		=	NS
č	3.	SHELL INSIDE DIA., inch		=	SID
č	4.	NUMBER OF BAFFLES		÷	NB
č	5.	BAFFLE SPACING, inch		=	BSP
č	6.	FOULING FACTOR		=	SFF
č	7.	INLET TEMPERATURE, OF		=	TS1
c	8.	OUTLET TEMPERATURE, OF		-	TS2
c	9.	SPECIFIC HEAT CAPACITY,	(Btu/lb oF)	=	SSHC
c	10.	VISCOSITY, CP	(BEU/10:01)		SVIS
c	11.	DENSITY, 1b/ft.^3		-	SDEN
c	12.	FLUID THERMAL CONDUCTIV	TTY (Btu/ft b o		SFK
c	12.	MASS FLOWRATE, 1b/hr.	, (Deu/re-M.O		SW
c	****** TO:	**************************************	******	-	
0					

PARAMETER (PI = 3.1415927, G=32.174)

CHARACTER*30 FTS,FSS CHARACTER TYPE

С

COMMON/TUBE1/ FTS, TYPE COMMON/TUBE2/ NT, TID, TOD, TP, TL, TNP

COMMON/TUBE3/ TFF, TT1, TT2, TSHC, TVIS, TDEN COMMON/TUBE4/ TFK, TW COMMON/TUBE5/ RET, PRT, TFRF, HT, TV COMMON/TUBE6/ TDELP, TRDELP, TOTDP COMMON/SHELL1/ FSS COMMON/SHELL2/ NS, SID, NB, BSP, SFF COMMON/SHELL3/ TS1, TS2, SSHC, SVIS, SDEN COMMON/SHELL4/ SFK, SW COMMON/SHELL5/ RES, PRS, SFRF, HS, SV COMMON/SHELL6/ SDELP COMMON/OUT/ Q, AREA, F, LMTD, U, Q1, CMTD REAL NT, NS, NB, LMTD OPEN (UNIT = 3, FILE='DATA82.DAT', STATUS='OLD', ERR= 18) OPEN (UNIT = 1, FILE='PRN') READ THE TUBE SIDE HEAT EXCHANGER PARAMETERS С С READ THE FLUID TYPE: FTS READ (3, 11, ERR=19)FTS FORMAT (A) 11 READ THE NUMBER OF TUBES: NT С С READ THE TUBE INSIDE DIAMETER, inch: TID READ THE TUBE OUTSIDE DIAMETER, inch: TOD С READ THE TUBE PITCH, inch: TP С с READ THE TYPE OF PITCH (TRIANGULAR: T) or (SQUARE: S): TYPE READ (3, *, ERR=19) NT, TID, TOD, TP READ (3, 12, ERR=19) TYPE 12 FORMAT (A) READ THE TUBE LENGTH, ft.: TL С С READ THE NUMBER OF PASSES: TNP č c READ THE TUBE SIDE FOULING FACTOR: TFF READ THE INLET TEMPERATURE, OF: TT1 READ THE OUTLET TEMPERATURE, OF: TT2 C C C READ THE SPECIFIC HEAT CAPACITY OF FLUID, (Btu/lb.of): TSHC READ THE FLUID VISCOSITY, CP: TVIS READ THE FLUID DENSITY, 1b/ft^3: TDEN READ THE FLUID THERMAL CONDUCTIVITY, Btu/hr.ft.of.: TFK С С с READ THE FLUID FLOWRATE, 1b/hr.: TW READ (3, *, ERR=19) TL, TNP, TFF, TT1 READ (3, *, ERR=19) TT2, TSHC, TVIS, TDEN READ (3, *, ERR=19) TFK, TW READ THE SHELL SIDE HEAT EXCHANGER PARAMETERS С READ THE FLUID TYPE: FSS С

READ (3, 13, ERR=19)FSS 13 FORMAT (A) READ THE NUMBER OF SHELL: NS С C READ THE SHELL SIDE INSIDE DIAMETER, inch: SID READ THE NUMBER OF BAFFLES: NB ¢ READ THE BAFFLE SPACING, inch: BSP С READ THE SHELL SIDE FOULING FACTOR, SFF READ THE INLET TEMPERATURE, OF: TS1 с С READ THE OULET TEMPERATURE, OF: TS2 С c c READ THE SHELL SIDE SPECIFIC HEAT CAPACITY, Btu/lb.oF.: SSHC READ THE FLUID VISCOSITY, CP: SVIS READ THE FLUID DENSITY, 1b/ft^3.: SDEN READ THE FLUID THERMAL CONDUCTIVITY, Btu/hr.ft.oF.: SFK C с С READ THE FLUID FLOWRATE, 1b/hr.: SW. READ (3, *, ERR=19) NS, SID, NB, BSP READ (3, *, ERR=19) SFF, TS1, TS2, SSHC READ (3, *, ERR=19) SVIS, SDEN, SFK, SW GO TO 15 18 WRITE (*, 20) FORMAT (3X, 'ERROR MESSAGE IN THE DATA VALUE') 20 GO TO 999 19 WRITE (*, 21) FORMAT (3X, 'DATA VALUE DOES NOT EXIST') 21 GO TO 999 с TUBE SIDE CALCULATION 15 CALL TUBE С SHELL SIDE CALCULATION CALL SHELL CALCULATE THE HEAT LOAD с Q = TW * TSHC * ABS(TT2 - TT1)c CALCULATE THE LOG MEAN TEMPERATURE DIFFERENCE (LMTD) VAL1 = (TS1-TT2)-(TS2-TT1)VAL2 = ALOG((TS1-TT2)/(TS2-TT1))LMTD = ABS(VAL1/VAL2) CALCULATE THE APPROACH FACTOR С R = (TS1-TS2)/(TT2-TT1)P = (TT2-TT1)/(TS1-TT1)IF (NS .EQ. 1.0 .AND. R .EQ. 1.0) THEN PX = P/(NS-NS*P+P)

```
SUM1 = (PX*(R*R+1.0)**0.5)/(1.0-PX)
        SUM2 = 2.0/PX-1.0-R+(R*R+1.0)**0.5
        SUM3 = 2.0/PX-1.0-R-(R*R+1.0)**0.5
        SUM4 = ALOG(SUM2/SUM3)
        F = SUM1/SUM4
        GO TO 100
        ELSEIF (NS .EQ. 1.0 .AND. R .GT. 1.0) THEN
        SUM5 = ((R*P-1.0)/(P-1.0))**(1.0/NS)
 55
        PX = (1.0-SUM5)/(R-SUM5)
        SUM6 = (R*R+1.0)**0.5/(R-1.0)
        SUM7 = ALOG((1.0-PX)/(1.0-R*PX))
        SUM41 = ((2.0/PX)-1.0-R+(R*R+1.0)**0.5)
        SUM42 = ((2.0/PX) - 1.0 - R - (R + 1.0) + 0.5)
        IF (SUM41 .LT. 0.0 .OR. SUM42 .LT. 0.0) THEN
        NS
            - NS+1.
        TNP = TNP+2.
        GO TO 55
        ELSE
        SUM43 = ALOG(SUM41/SUM42)
        ENDIF
        F \approx (SUM6 \times SUM7) / SUM43
        IF (F .LT. 0.8) THEN
        NS = NS+1.0
        TNP = TNP+2.
        GO TO 55
        ELSE
        END IF
        GO TO 100
        ELSEIF (NS .GE. 2.0 .AND. NT .GE. 4.0) THEN
        SUM8 = (R*R+1.0)**0.5/(2.0*(R-1.0))
        SUM9 = ALOG((1.0-P)/(1.0-P*R))
        SUM10 = 2/P-1.0-R+((2/P)*((1.0-P)*(1-P*R))**0.5)+(R*R+1.)**0.5
        SUM11 = 2/P-1.0-R+((2/P)*((1.0-P)*(1.-P*R))**0.5)-(R*R+1.)**0.5
        F = (SUM8 * SUM9) / ALOG(SUM10 / SUM11)
        ENDIF
с
        CORRECTED MEAN TEMPERATURE DIFFERENCE
100
        CMTD = F*LMTD
        CALCULATE THE OVERALL HEAT TRANSFER COEFFICIENT
C
С
        Btu/hr.ft^2.oF
        U = 1.0/(1./HT+1./HS+TFF+SFF)
Ċ
        CALCULATE THE OUTSIDE AREA OF THE HEAT EXCHANGER UNIT ft^2.
        AREA = (TOD/12.0*PI*NT*NS*TL)
```

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688
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```
Q1 = AREA*U*CMTD
CALL RESULT
C FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
WRITE (1, *) CHAR(12)
CLOSE (UNIT =3, STATUS='KEEP')
CLOSE (UNIT = 1)
999 STOP
END
```

С ***** c c *THIS PROGRAM CALCULATES THE TUBE SIDE HEAT TRANSFER COEFFICIENT, *REYNOLDS NUMBER, PRANDTL NUMBER, FRICTION FACTOR AND PRESSURE DROP.* С SUBROUTINE TUBE PARAMETER (PI = 3.1415927, G=32.174) CHARACTER*30 FTS, FSS CHARACTER TYPE COMMON/TUBE1/ FTS, TYPE COMMON/TUBE2/ NT, TID, TOD, TP, TL, TNP COMMON/TUBE3/ TFF, TT1, TT2, TSHC, TVIS, TDEN COMMON/TUBE4/ TFK, TW COMMON/TUBE5/ RET, PRT, TFRF, HT, TV COMMON/TUBE6/ TDELP, TRDELP, TOTDP COMMON/SHELL1/ FSS COMMON/SHELL1/ FSS COMMON/SHELL2/ NS, SID, NB, BSP, SFF COMMON/SHELL3/ TS1, TS2, SSHC, SVIS, SDEN COMMON/SHELL4/ SFK, SW COMMON/SHELL5/ RES, PRS, SFRF, HS, SV COMMON/SHELL5/ SDELP COMMON/OUT/ Q, AREA, F, LMTD, U, Q1, CMTD REAL NT, NS, NB, LMTD С CALCULATE THE FLOW AREA: TFA, ft^2 TFA = (PI*TID**2*NT)/(4.0*144.0*TNP)С CALCULATE THE MASS VELOCITY, 1b/hr.ft^2 GT = TW/TFAC CALCULATE THE REYNOLDS NUMBER: RET RET = (GT*TID)/(12.0*2.42*TVIS) С CALCULATE THE PRANDTL NUMBER: PRT PRT = (TSHC*TVIS*2.42)/TFK с с CHECK THAF THE REYNOLDS NUMBER IS IN THE TRANSITION REGION (i.e. 2000<RET<10,000) IF (RET .GT, 2000.0 .AND. RET .LT. 10000.0) THEN X1 = RET/1000.0 $X2 = 10.0 \star TID/TL$ X3 = (1.0 - (X1/10.0) * * 0.256)X4 = X2 * X3B1 = (-3.08+3.075*X1+(0.32567*X1**2)-(0.02185*X1**3))*X4HT = (16.1/TOD)*(B1*TFK*PRT**0.33)GO TO 110

ELSE IF (RET .GT. 10000.0) THEN

```
HT = 0.027*TFK*(12.0/TID)*(RET**0.8)*(PRT**0.33)
        ENDIF
        HT = HT*TID/TOD
 110
        CALCULATE THE TUBE FRICTION FACTOR
С
        IF (RET .GT. 1000.0) THEN
        TFRF = 0.0029/(RET**0.25603)
        ELSE
        ENDIF
        CALCULATE THE FLUID VELOCITY, ft./s.
С
        TV = GT/(3600.0 * TDEN)
        CALCULATE THE TUBE SIDE PRESSURE DROP, psi.
с
        TDELP = (TFRF*(GT/3600.0)**2*TL*TNP)/(2.0*G*TDEN*TID/12.0)
        CALCULATE THE PRESSURE DROP OF RETURN LOSSES, psi.
С
        USING FRANK'S CORRELATION.
С
        TRDELP = (2.5*TNP*TDEN*TV**2)/(2*G*144.0)
c
c
        CALCULATE THE TOTAL PRESSURE DROP EXCLUDING THE NOZZLE IN THE
        TUBE SIDE, psi.
        TOTDP = TDELP+TRDELP
        RETURN
        END
```

С ****** 0000 *THIS PROGRAM CALCULATES THE SHELL SIDE HEAT TRANSFER *COEFFICIENT, REYNOLDS NUMBER, PRANDTL NUMBER, FRICTION FACTOR * *AND PRESSURE DROP. ***** SUBROUTINE SHELL PARAMETER (PI = 3.1415927, G=32.174) CHARACTER*30 FTS, FSS CHARACTER TYPE COMMON/TUBE1/ FTS, TYPE COMMON/TUBE2/ NT, TID, TOD, TP, TL, TNP COMMON/TUBE3/ TFF, TT1, TT2, TSHC, TVIS, TDEN COMMON/TUBES/ TFK, TW COMMON/TUBES/ RET, PRT, TFRF, HT, TV COMMON/TUBES/ RET, PRT, TFRF, HT, TV COMMON/SHELL1/ FSS COMMON/SHELL2/ NS, SID, NB, BSP, SFF COMMON/SHELL3/ TS1, TS2, SSHC, SVIS, SDEN COMMON/SHELL3/ TS1, TS2, SSHC, SVIS, SDEN COMMON/SHELL4/ SFK, SW COMMON/SHELL6/ SDELP COMMON/OUT/ Q, AREA, F, LMTD, U, Q1, CMTD REAL NT, NS, NB, LMTD С CHECK WHETHER PITCH TYPE IS TRIANGULAR OR SQUARE IF (TYPE .EQ. 'T' .OR. TYPE .EQ. 't') THEN DE = ((1.72*TP**2)-(0.5*PI*TOD**2))/(0.5*PI*12.0*TOD) GO TO 120 ELSE IF (TYPE .EQ. 'S' .OR. TYPE .EQ. 's') THEN DE = 4.*((TP**2)-(PI*TOD**2/4.0))/(PI*12.0*TOD)ENDIF CALCULATE THE TUBE CLEARANCE, inch С TC = TP - TOD120 С CALCULATE THE SHELL FLOW AREA, ft^2. IF (NS .GE. 2.0 .AND. NT .GE. 4.0) THEN SFA = (0.5*(SID*TC*BSP))/(144.0*TP) GO TO 130 ELSE SFA = (SID*TC*BSP)/(144.0*TP)ENDIF

с	CALCULATE THE SHELL SUPERFICIAL FLOW RATE, 1b/ft^2.hr.
130	GS = SW/SFA
С	CALCULATE THE SHELL VELOCITY, ft./sec.
	SV = SW/(3600.0*SDEN*SFA)
с	CALCULATE THE REYNOLDS NUMBER
	RES = (DE*GS)/(2.42*SVIS)
С	CALCULATE THE PRANDTL NUMBER
	PRS = (2.42*SSHC*SVIS)/SFK
С	CALCULATE THE SHELL SIDE HEAT TRANSFER COEFFICIENT
	HS = 0.36/DE*SFK*(RES**0.55)*(PRS**0.333)
C C	CHECK WHETHER THE SHELL'S REYNOLDS NUMBER IS LESS OR GREATER THAN 500
	IF (RES .LT. 500.0) THEN
	SFRF = 0.11183/(RES**0.59246)
	GO TO 140
	ELSEIF (RES .GT. 500.0) THEN
	SFRF = 0.01159/(RES**0.18597) ENDIF
с	CALCULATE THE SHELL SIDE PRESSURE DROP, psi.
140	VAL1 = SFRF*(GS/3600.0)**2*SID/12.0*(NB+1.) VAL2 = 64.4*SDEN*DE SDELP = (VAL1/VAL2)*NS
с	RETURN

RETURN END С ċ THIS PROGRAM PRINTS THE RESULTS OF THE SHELL AND TUBE HEAT С EXCHANGER ONTO A PRINTER С ********** SUBROUTINE RESULT CHARACTER*30 FTS, FSS CHARACTER TYPE COMMON/TUBE1/ FTS, TYPE COMMON/TUBE2/ NT, TID, TOD, TP, TL, TNP COMMON/TUBE3/ TFF, TT1, TT2, TSHC, TVIS, TDEN COMMON/TUBE4/ TFK, TW COMMON/TUBE5/ RET, PRT, TFRF, HT, TV COMMON/TUBEG/ TDELP, TRDELP, TOTDP COMMON/SHELL1/ FSS COMMON/SHELL2/ NS, SID, NB, BSP, SFF COMMON/SHELL3/ TS1, TS2, SSHC, SVIS, SDEN COMMON/SHELL3/ TS1, TS2, SSHC, SVIS, SDEN COMMON/SHELL5/ RES, PRS, SFRF, HS, SV COMMON/SHELL6/ SDELP COMMON/OUT/ Q, AREA, F, LMTD, U, Q1, CMTD REAL NT, NS, NB, LMTD WRITE (1, 150) 150 FORMAT (///,18X,'*SHELL AND TUBE HEAT EXCHANGER RATING*') WRITE (1, 160) 160 FORMAT (78(1H*)) WRITE (1, 170) 170 FORMAT (40X, 'TUBE SIDE', 10X, 'SHELL SIDE') WRITE (1, 180) 180 FORMAT (78(1H-)) WRITE (1, 190) FTS, FSS 190 FORMAT (2X, 'FLUID TYPE: ', T40, A15, T60, A15) WRITE (1, 200) TW, SW 200 FORMAT (2X, 'FLUID FLOW RATE , 1b/hr.:', T40, F10.0, T60, F10.0) WRITE (1, 210) TSHC, SSHC 210 FORMAT (2X, 'SPECIFIC HEAT CAPACITY, Btu/lb.oF:', T40, F8.3, T60, F8.3) WRITE (1, 220)TDEN, SDEN FORMAT (2X, 'FLUID DENSITY, 1b/ft^3:',T40,F8.3,T60,F8.3) 220 WRITE (1, 230) TVIS, SVIS 230 FORMAT (2X, 'FLUID VISCOSITY, CP:', T40, F8.3, T60, F8.3) WRITE (1, 240) TFK, SFK 240 FORMAT (2X, 'FLUID THERMAL COND. Btu/ft.h.oF:', T40, F8.3, T60, F8.3) WRITE (1, 250) TT1, TS1 250 FORMAT (2X, 'FLUID INLET TEMPERATURE, oF: ', T40, F8.0, T60, F8.0)

260	WRITE (1, 260) TT2, TS2 FORMAT (2X,'FLUID OUTLET TEMPERATURE, oF:',T40,F8.0,T60,F8.0)
270	WRITE (1, 270) TID, SID FORMAT (2X,'INSIDE DIAMETER, inch:',T40,F8.3,T60,F8.3)
280	WRITE (1, 280) TOD FORMAT (2X,'OUTSIDE DIAMETER, inch:',T40,F8.3)
	IF (TYPE .EQ. 'T' .OR. TYPE .EQ. 't') THEN
290	WRITE (1, 290) TP FORMAT (2X,'TUBE PITCH, inch:',T40,F8.3,T60,'TRIANGULAR PITCH') ELSEIF (TYPE .EQ. 'S' .OR. TYPE .EQ. 'S') THEN
300	WRITE (1, 300) TP FORMAT (2X,'TUBE PITCH, inch:',T40,F8.3,T60,'SQUARE PITCH') ENDIF
310	WRITE (1, 310) TL FORMAT (2X,'TUBE LENGTH, ft.:',T40,F8.3)
315	WRITE (1, 315) NT FORMAT (2X,'NUMBER OF TUBES:', T40, F6.0)
320	WRITE (1, 320) TNP, NS FORMAT (2X,'NUMBER OF PASSES:',T40,F4.0,T60,F4.0)
340	WRITE (1, 340) NB FORMAT (2X,'NUMBER OF BAFFLES:',T60,F4.0)
350	WRITE (1, 350) BSP FORMAT (2X,'BAFFLE SPACING, inch:',T60,F4.1)
360	WRITE (1, 360) TV, SV FORMAT (2X,'FLUID VELOCITY, ft/sec.:',T40,F8.3,T60,F8.3)
370	WRITE (1, 370) RET, RES FORMAT (2X,'REYNOLDS NUMBER:',T40,F9.0,T60,F9.0)
380	WRITE (1, 380) PRT, PRS FORMAT (2X,'PRANDTL NUMBER:',T40,F8.4,T60,F8.4)
390	WRITE (1, 390) TFRF, SFRF FORMAT (2X,'FRICTION FACTOR (ft^2/in^2):',T40,F8.5,T60,F8.5)
400	WRITE (1, 400) HT, HS FORMAT (2X,'HEAT COEFF., Btu/hr.ft^2.oF:',T40,F9.1,T60,F9.1)
4 10	WRITE (1, 410) TOTDP, SDELP FORMAT (2X,'PRESSURE DROP, psi:',T40,F8.4,T60,F8.4)
420	WRITE (1, 420) TFF, SFF FORMAT (2X,'FOULING FACTOR:',T40,F8.4,T60,F8.4)
430	WRITE (1, 430) F FORMAT (2X,'APPROACH FACTOR, F:', T40, F8.4)

440	WRITE (1, 440) LMTD FORMAT (2X,'LOG MEAN TEMP. DIFF. oF:',T40,F8.2)
450	WRITE (1, 450) CMTD FORMAT (2X,'CORRECTED MEAN TEMP. DIFF. oF:',T40,F8.2)
460	WRITE (1, 460) U FORMAT (2X,'OVERALL HEAT COEFF., Btu/hr.ft^2.oF:',T40,F9.1)
470	WRITE (1, 470) AREA FORMAT (2X,'OUTSIDE AREA OF UNIT, ft^2:',T40,F9.1)
480	WRITE (1, 480) Q FORMAT (2X,'HEAT LOAD ON UNIT, Btu/hr:',T40,F12.0)
490	WRITE (1, 490) Q1 FORMAT (2X,'HEAT TRANSFERRED, Btu/hr:',T40,F12.0)
500	WRITE (1, 500) FORMAT (1H ,78(1H-))
с	FORM FEED THE PRINTING PAPER TO THE TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR(12)
	RETURN END

PROGRAM PROG83

с	********	
c	THIS PROGRAM RATES DOUBLE PIPE HEAT EXCHANGERS USING	•
c	BARE TUBES OR LONGITUDINAL FINNED TUBES.	
c	THE PROGRAM CALCULATES THE HEAT LOAD, THE LOG MEAN TEMPERATURE	
c		
	DIFFERENCE FOR COUNTER CURRENT FLOW, THE CORRECTED FILM HEAT TRANSFE	5 R
С	COEFFICIENTS IN BOTH THE TUBE AND THE SHELL SIDES, THE OVERALL HEAT	
с	TRANSFER COEFFICIENT AND THE HEAT TRANSFER SURFACE. THE PROGRAM	
с	FURTHER CALCULATES THE FRICTION FACTOR (Using correlations given by	
с	Dennis Wright: Basic Programs for chemical engineers, Van Nostrand	
С	Reinhold Company, NY 1986) AND THE PRESSURE DROP IN THE TUBE AND THE	3
С	SHELL SIDES OF THE DOUBLE PIPE HEAT EXCHANGERS.	
С	***************************************	*
С	HI = HEAT-TRANSFER FILM COEFFICIENT, Btu/ft^2 of.h	
с	K = FLUID THERMAL CONDUCTIVITY, Btu/ft.h.oF	
С	RE = REYNOLDS NUMBER	
С	PR = PRANDTL NUMBER	
С	DEQ = EQUIVALENT DIAMETER, ft.	
с	VIS = FLUID VISCOSITY (FLOWING), cP.	
с	VISW = FLUID VISCOSITY (AT WALL), cP.	
с	VIST = FLUID VISCOSITY (TUBE SIDE), CP.	
С	VISS = FLUID VISCOSITY (SHELL SIDE - bare tube), cP.	
С	VISF = FLUID VISCOSITY (SHELL SIDE - finned tube), cP.	
с	J = HEAT-TRANSFER FACTOR	
с	$G = MASS FLOW RATE, lb/h.ft^2.$	
c	CP = FLUID SPECIFIC HEAT, Btu/lb.oF.	
ċ	AREA = HEAT TRANSFER SURFACE AREA, ft^2 .	
č	IDT = INSIDE-TUBE DIAMETER, in.	
č	IDF = INSIDE DIAMETER (finned tube), in.	
č	FOL = FOULING FACTOR	
č	HIF = HEAT-TRANSFER FILM COEFFICIENT CORRECTED	
č	FOR FOULING, Btu/ft^2.oF.h.	
č	IDS = INSIDE SHELL DIAMETER, in.	
č	ODT = OUTSIDE TUBE DIAMETER, in.	
c	DELTW = TUBE WALL THICKNESS, ft.	
c	KT = THERMAL CONDUCTIVITY OF TUBE MATERIAL, Btu/ft.h.of.	
c	AI = INSIDE HEAT-TRANSFER AREA-TUBE, ft^2/ft .	
c	AO = OUTSIDE HEAT-TRANSFER AREA-TUBE, ft $2/ft$ (FOR FINNED	
c	TUBES INCLUDES FIN AREA).	
c	UO = OVERALL HEAT-TRANSFER COEFFICIENT, Btu/h.ft^2.oF.	
č	NFA = NET FREE CROSS-SECTIONAL AREA, SHELL SIDE, ft^2 .	
č	$AF = FINNED TRANSFER AREA, ft^2.$	
č	X = PARAMETER USED IN FIN EFFICIENCY CALCULATION	
č	FE = FIN EFFICIENCY (%).	
č	ESEF = EFFECTIVE SURFACE EFFICIENCY FOR FINS (%).	
c	HIFD = CORRECTED FILM HEAT-TRANSFER RATE, $Btu/ft^2.h.oF$	
c	TDEN = TUBE SIDE FLUID DENSITY, $1b/ft^3$.	
c	SDEN = SHELL SIDE FLUID DENSITY, 1b/ft^3.	
c	NTP = NUMBER OF TUBE PASSES.	
c	NIP = NUMBER OF IUDE PASSES. NSP = NUMBER OF SHELL PASSES (bare-tube).	
c	NFP = NUMBER OF SHELL PASSES (finned-tube).	
c	FOLT = FOULING FACTOR OF THE TUBE SIDE.	
c	FOLS = FOULING FACTOR OF THE SHELL SIDE.	
c	FOLF = FOULING FACTOR OF THE SHELL SIDE (Dare-tube).	
c		
c		
	FTSHELL = FLUID NAME IN THE SHELL SIDE	
с	FLUIDT = FLUID PHASE FLOW IN THE TUBE SIDE.	

```
С
         FLUIDS = FLUID PHASE FLOW IN THE SHELL SIDE.
                   = CHECKS TO DETERMINE WHETHER THE DOUBLE PIPE HEAT
00000000
         TUBE
                      EXCHANGER IS BARE-TUBED OR FINNED-TUBE.
         LMTD
                   = LOG MEAN TEMPERATURE DIFFERENCE (oF).
                   = TUBE LENGTH, ft.
         TL
                   = FLUID FLOWRATE TUBE SIDE, lb/hr.
= FLUID FLOWRATE SHELL SIDE (BARE-TUBE), lb/hr.
         WΤ
         ws
                   = FLUID FLOWRATE SHELL SIDE (FINNED-TUBE), 1b/hr.
          WF
c
          ********
          CHARACTER*30 FLUIDT, FLUIDS, FLUIDF, TUBE, FTTUBE, FTSHELL
         EXTERNAL HTFCV, HTFCL, HTFCJ
          REAL IDTF, IDSF, IDFF, ODTF, ODSF, ODFF
         REAL IDT, ODT, KT
         REAL IDS, ODS, KS, KST
REAL IDF, ODF, KF, KFT
          REAL LMTD
         COMMON TUBE
          COMMON/DATA/FTTUBE, FTSHELL, TDEN, SDEN
COMMON/DATA1/FLUIDT,IDT,ODT,WT,CPT,VIST,KT,TT1,TT2,FOLT,TL,NTP
         COMMON/DATA2/FLUIDS, IDS, ODS, WS, CPS, VISS, KS, TS1, TS2, FOLS, KST, NSP
COMMON/DATA3/FLUIDF, IDF, ODF, WF, CPF, VISF, KF, TF1, TF2, FOLF,
      * FHT,FTHICK,NF,KFT,NFP
COMMON/DATA4/ PI
COMMON/DATA5/ IDTF, IDSF, IDFF, ODTF, ODSF, ODFF
          COMMON/RES1/VELT, RET, PRT, FRICTT, DROPT, HTFT, HIFTU
          COMMON/RES2/VELS, RES, PRS, FRICTS, DROPS, HTFS, HIFS
          COMMON/RES3/VELF, REF, PRF, FRICTF, DROPF, HTFF, HIFF, X, FE, ESEF, HIFD,
         UO
          COMMON/RES4/ LMTD, AREA, Q
         OPEN (UNIT=3, FILE='DATA83.DAT', STATUS='OLD', ERR=18)
         OPEN (UNIT= 1, FILE='PRN')
         PI = 3.1415927
С
         TUBE-SIDE HEAT-TRANSFER CALCULATIONS
С
          INPUT NAME OF FLUID IN THE TUBE SIDE
         READ (3, 9, ERR=19) FTTUBE
 9
         FORMAT (A)
         INPUT FLUID PHASE FLOW (e.g. liquid/vapor)
С
         READ (3, 11, ERR=19) FLUIDT
 11
         FORMAT (A)
         READ THE INSIDE TUBE DIAMETER, inch.: IDT
С
         READ THE OUTSIDE TUBE DIAMETER, inch.: ODT
READ THE FLUID FLOWRATE IN THE TUBE SIDE, lb/hr.: WT
с
С
         READ THE FLUID SPECIFIC HEAT CAPACITY, Btu/lb.oF.: CPT
С
С
         READ THE FLUID VISCOSITY, CP: VIST
```

READ THE FLUID THERMAL CONDUCTIVITY, KT: Btu/hr.ft.oF: KT

698

С

READ THE FLUID INLET TEMPERATURE, oF: TT1 READ THE FLUID OUTLET TEMPERATURE, oF: TT2 READ THE TUBE SIDE FOULING FACTOR: FOLT с č č С READ THE TUBE SIDE FLUID DENSITY, 1b/ft^3: TDEN С READ THE TUBE LENGTH, ft.: TL с READ THE NUMBER OF TUBE PASSES: NTP READ (3, *, ERR=19) IDT, ODT, WT READ (3, *, ERR=19) CPT, VIST, KT READ (3, *, ERR=19) TT1, TT2, FOLT READ (3, *, ERR=19) TDEN, TL, NTP GO TO 10 CHECK WHETHER THE HEAT EXCHANGER IS BARE-TUBE OR FINNED-TUBE С С i.e., WHETHER THE HEAT EXCHANGER HAS FINS С INPUT NAME OF FLUID IN THE SHELL SIDE 5 READ (3, 14, ERR=19) FTSHELL FORMAT (A) 14 READ (3, 22, ERR=19) TUBE 22 FORMAT (A) IF (TUBE .EQ. 'FINS' .OR. TUBE .EQ. 'fins') THEN GO TO 33 ELSEIF (TUBE .EQ. 'BARE-TUBE'.OR. TUBE .EQ. 'bare-tube') THEN С READ DATA VALUES FOR THE SHELL-SIDE BARE-TUBE с INPUT FLUID PHASE FLOW (e.g. liquid/vapor) READ (3, 34, ERR=19) FLUIDS 34 FORMAT(A) С READ THE INSIDE DIAMETER, inch.: IDS READ THE OUTSIDE DIAMETER, inch.: ODS READ THE FLUID FLOWRATE, lb/hr.: WS READ THE FLUID SPECIFIC HEAT CAPACITY, Btu/lb.oF: CPS READ THE FLUID VISCOSITY, CP.: VISS READ THE FLUID THERMAL CONDUCTIVITY, Btu/ft.hr.oF: KS READ THE FLUID INLET TEMPERATURE, OF: TS1 READ THE FLUID OUTLET TEMPERATURE, OF: TS2 READ THE SHELL SIDE FOULING FACTOR: FOLS READ THE TUBE THERMAL CONDUCTIVITY, Btu/ft.hr.oF: KST READ THE FLUID DENSITY, 1b/ft^3.: SDEN READ THE NUMBER OF SHELL PASSES: NSP READ (3, *, ERR=19) IDS, ODS, WS READ (3, *, ERR=19) CPS, VISS, KS READ $(3, \star, ERR=19)$ TS1, TS2, FOLS READ $(3, \star, ERR=19)$ KST, SDEN, NSP

ENDIF

	ENDIF
с	CALCULATE THE LOG MEAN TEMPERATURE DIFFERENCE, LMTD (oF)
	CALL LOGMTD (TS1, TS2, TT1, TT2, LMTD)
	GO TO 20
с	READ DATA VALUES FOR THE SHELL-SIDE FINNED TUBE
с	INPUT PHASE FLOW OF FLUID IN THE SHELL SIDE (e.g. Liquid/vapor)
33 44	READ (3, 44, ERR=19) FLUIDF FORMAT (A)
000000000000000	READ THE NUMBER OF FINS: NF READ THE INSIDE DIAMETER, inch.: IDF READ THE OUTSIDE DIAMETER, inch.: ODF READ THE FLUID SIDE DIAMETER, inch.: ODF READ THE FLUID SPECIFIC HEAT, Btu/lb.oF: CPF READ THE FLUID VISCOSITY, CP.: VISF READ THE FLUID UISCOSITY, CP.: VISF READ THE FLUID UISCOSITY, CP.: VISF READ THE FLUID INLET TEMPERATURE, OF: TF1 READ THE FLUID OUTLET TEMPERATURE, OF: TF2 READ THE FLUID OUTLET TEMPERATURE, OF: TF2 READ THE FLUID OUTLET TEMPERATURE, OF: TF2 READ THE FINS HEIGHT, inch.: FHT READ THE FINS HEIGHT, inch.: FTHICK READ THE FINS THICHNESS, inch.: FTHICK READ THE FINS THICHNESS, inch.: SDEN READ THE FUES SIDE THERMAL CONDUCTIVITY, Btu/hr.ft.oF: KFT READ THE FUES SIDE THERMAL CONDUCTIVITY, Btu/hr.ft.oF: KFT READ THE NUMBER OF PASSES: NFP READ (3, *, ERR=19) NF READ (3, *, ERR=19) CFP, VISF, KF READ (3, *, ERR=19) TF1, TF2, FOLF READ (3, *, ERR=19) TF1, TF2, FOLF READ (3, *, ERR=19) KFT, SDEN, NFF
с	CALCULATE THE LOG MEAN TEMPERATURE DIFFERENCE, LMTD (oF)
-	CALL LOGMTD (TF1, TF2, TT1, TT2, LMTD)
с	CALCULATE THE HEAT LOAD ON UNIT, Btu/hr.
	Q = WF*CPF*(TF1-TF2)
с	SHELL-SIDE CALCULATIONS (FINNED-TUBE)
	GO TO 30
18 100	WRITE (*, 100) FORMAT(//,6X,'DATA FILE DOES NOT EXIST') GO TO 999
19	WRITE (*, 110)

110	FORMAT(//,6X,'ERROR MESSAGE IN THE DATA VALUE') GO TO 999
с	TUBE-SIDE CALCULATIONS
10	CALL PROG1 GO TO 5
с	SHELL-SIDE CALCULATIONS (BARE-TUBE)
20	CALL PROG2
	GO TO 40
с	SHELL-SIDE CALCULATIONS (FINNED-TUBE)
30	CALL PROG3
с	CALCULATE THE HEAT TRANSFER SURFACE AREA, (AREA, ft^2).
40	AREA = Q/(UO*LMTD)
	CALL RESULTS
	CLOSE (3, STATUS='KEEP') CLOSE (1)
999	STOP END
с с с с	**************************************
	SUBROUTINE PROG1
	CHARACTER*30 FLUIDT, FTTUBE, FTSHELL,FLUIDF
	REAL IDT, ODT, KT REAL IDTF, IDSF, IDFF, ODTF, ODSF, ODFF REAL IDF, ODF, KF, KFT
*	COMMON/DATA/FTTUBE, FTSHELL, TDEN, SDEN COMMON/DATA1/FLUIDT,IDT,ODT,WT,CPT,VIST,KT,TT1,TT2,FOLT,TL,NTP COMMON/DATA3/FLUIDF,IDF,ODF,WF,CPF,VISF,KF,TF1,TF2,FOLF, FHT,FTHICK,NF,KFT,NFP COMMON/DATA4/FI COMMON/DATA5/IDTF, IDSF, IDFF, ODTF, ODSF, ODFF
	COMMON/RES1/VELT, RET, PRT, FRICTT, DROPT, HTFT, HIFTU
	IDTF = IDT/12.0 ODTF = ODT/12.0 DEQT = IDTF

C CALCULATE THE CROSS-SECTIONAL AREA OF TUBE

AT = (PI*IDTF**2)/4.0

C CALCULATE THE MASS VELOCITY, lb/ft^2.hr.

GT = WT/AT

C CALCULATE THE FLUID VELOCITY, ft./sec.

VELT = GT/(3600.0*TDEN)

C CALCULATE THE REYNOLDS NUMBER

RET = (GT*DEQT)/(2.42*VIST)

C CALCULATE THE PRANDTL NUMBER

PRT = (CPT*2.42*VIST)/KT

C CALCULATE THE FRICTION FACTOR (Dennis Wright: Basic Programs C for chemical engineers: Van Nostrand Reinhold Company, NY 1986).

IF (RET .GT. 2100.0) THEN

FRICTT = 0.00033/(RET**0.2)
ELSEIF (RET .LT. 2100.0) THEN
FRICTT = 16/RET
ENDIF

C CALCULATE THE PRESSURE DROP

DROPT = (4.0*FRICTT*TL*NTP*(GT/3600.0)**2)/(64.4*TDEN*DEQT)

C CHECK WHETHER FLUID IN THE TUBE SIDE IS LIQUID OR VAPOR.

IF (FLUIDT .EQ. 'LIQUID'.OR. FLUIDT .EQ. 'liquid') THEN GO TO 30 ELSEIF (FLUIDT .EQ. 'VAPOR' .OR. FLUIDT .EQ. 'vapor') THEN

C CALCULATE THE HEAT-TRANSFER FILM COEFFICIENT IN THE VAPOR STATE

HTFT = HTFCV (CPT,GT,DEQT) ENDIF GO TO 90

- 30 IF(RET .LT. 2100.0) THEN
 GO TO 50
 ELSE
 HTFT = HTFCL (RET,PRT,KT,DEQT)
 ENDIF
 GO TO 90
- 50 HTFT = HTFCJ (RET, PRT, KT, DEQT, TL)

C C	CALCULATE THE HEAT-TRANSFER FILM COEFFICIENT CORRECTED FOR FOULING, Btu/hr.ft^2.oF
90	HIFTU = HTFT/(1.0+HTFT*FOLT)
	RETURN END
с с с с с с с	**************************************
	SUBROUTINE PROG2
	CHARACTER*30 FLUIDS, FLUIDT, FTSHELL,FTTUBE REAL IDS, IDT, ODS, ODT, KS, KT REAL IDTF, IDSF, IDFF, ODTF, ODSF, ODFF REAL LMTD
	COMMON/DATA/FTTUBE, FTSHELL, TDEN, SDEN COMMON/DATA1/FLUIDT,IDT,ODT,WT,CPT,VIST,KT,TT1,TT2,FOLT,TL,NTP COMMON/DATA2/FLUIDS,IDS,ODS,WS,CPS,VISS,KS,TS1,TS2,FOLS,KST,NSP COMMON/DATA4/PI COMMON/DATA5/IDTF, IDSF, IDFF, ODTF, ODSF, ODFF
	COMMON/RES1/VELT,RET,PRT,FRICTT,DROPT,HTFT,HIFTU COMMON/RES2/VELS,RES,PRS,FRICTS,DROPS,HTFS,HIFS
	COMMON/RES4/LMTD, AREA, Q
	IDSF = IDS/12.0 ODSF = ODS/12.0 DEQS = IDSF-ODTF RATIO1 = IDT/ODT
с	CALCULATE THE CROSS-SECTIONAL AREA
	AS = (PI*IDSF**2)/4.0
с	CALCULATE THE MASS VELOCITY, Ib/ft^2.hr.
	GS = WS/AS
с	CALCULATE THE FLUID VELOCITY, ft./sec.
	VELS = GS/(3600.0*SDEN)
с	CALCULATE THE REYNOLDS NUMBER
	RES = (GS*DEQS)/(2.42*VISS)
с	CALCULATE THE PRANDTL NUMBER
	PRS = (CPS*2.42*VISS)/KS

C CALCULATE THE FRICTION FACTOR (Dennis Wright: Basic Programs for C chemical engineers, Van Nostrand Reinhold Company, NY 1986.)

С

CALCULATE THE FRICTION FACTOR

CALL FFACTOR (RES, FRICTS)

с CALCULATE THE PRESSURE DROP, psi. CALL PDROP (FRICTS, TL, NSP, GS, SDEN, DEQS, DROPS) IF (FLUIDS .EQ. 'LIQUID' .OR. FLUIDS .EQ. 'liquid') THEN GO TO 30 ELSEIF (FLUIDS .EQ. 'VAPOR' .OR. FLUIDS .EQ. 'vapor') THEN CALCULATE THE HEAT TRANSFER FILM COEFFICIENT IN THE VAPOR STATE с HTFS = HTFCV(CPS,GS,DEQS) ENDIF GO TO 90 IF (RES .LT. 2100.0) THEN GO TO 50 30 ELSE HTFS = HTFCL (RES, PRS, KS, DEQS) ENDIF GO TO 90 50 HTFS = HTFCJ (RES, PRS, KS, DEQS, TL) CALCULATE THE HEAT-TRANSFER FILM COEFFICIENT CORRECTED c c FOR FOULING. 90 HIFS = HTFS/(1.0+HTFS*FOLS) с CALCULATE THE OVERALL HEAT TRANSFER COEFFICIENT. DELTW = (ODT/12.) - (IDT/12.0)RATIO2 = (DELTW/KST)+(1.0/(HIFTU*RATIO1))+(1.0/HIFS) UO = 1.0/RATIO2RETURN END CCCCC PROGRAM CALCULATES HEAT-TRANSFER FILM COEFFICIENT IN THE SHELL SIDE (FINNED-TUBE) AND THE OVERALL HEAT TRANSFER COEFFICIENT OF THE HEAT EXCHANGER. SUBROUTINE PROG3 CHARACTER*30 FLUIDT, FLUIDF, FTSHELL, FTTUBE

REAL IDF, ODF, KF, KFT, KT, NFA, IDT, ODT

```
REAL IDTF, IDSF, IDFF, ODTF, ODSF, ODFF
         COMMON/DATA/FTTUBE, FTSHELL, TDEN, SDEN
COMMON/DATA1/FLUIDT,IDT,ODT,WT,CPT,VIST,KT,TT1,TT2,FOLT,TL,NTP
COMMON/DATA3/FLUIDF,IDF,ODF,WF,CPF,VISF,KF,TF1,TF2,FOLF,
        FHT, FTHICK, NF, KFT, NFP
         COMMON/DATA4/PI
         COMMON/DATA5/IDTF, IDSF, IDFF, ODTF, ODSF, ODFF
         COMMON/RES3/VELF, REF, PRF, FRICTF, DROPF, HTFF, HIFF, X, FE, ESEF, HIFD,
     * UO
         IDFF = IDF/12.0
         ODFF = ODF/12.0
         ODTF = ODT/12.0
         AF = (2.0 \star FHT \star FLOAT(NF))/12.0
         AO = (PI*ODTF)/12.0+AF
с
         CALCULATE CROSS-SECTIONAL AREA, SHELL SIDE WITHOUT FINS, ft^2.
         CSA = (PI*(IDFF**2-ODTF**2))/4.0
         CALCULATE NET FREE CROSS-SECTIONAL AREA, SHELL SIDE, ft^2.
С
         NFA = CSA-(FHT*FTHICK*FLOAT(NF))/144.0
С
         CALCULATE THE CROSS-SECTIONAL AREA.
         CAF = (PI*IDFF**2)/4.0
         CALCULATE THE MASS VELOCITY, lb/ft^2.hr.
С
         GF = WF/NFA
         CALCULATE THE FLUID VELOCITY, ft./sec.
С
         VELF = GF/(3600.0*SDEN)
         CALCULATE THE WETTED PERIMETER.
С
         DENOM = (PI*(IDFF+ODTF))~(FTHICK*FLOAT(NF))/12.0+AF
         DEQF = (4.0 * NFA) / DENOM
с
         CALCULATE THE REYNOLDS NUMBER
         REF = (GF*DEQF)/(2.42*VISF)
с
         CALCULATE THE PRANDTL NUMBER
         PRF = (CPF*2.42*VISF)/KF
         CALCULATE THE FRICTION FACTOR
С
         CALL FFACTOR (REF, FRICTF)
         CALCULATE THE PRESSURE DROP
         CALL PDROP (FRICTF, TL, NFP, GF, SDEN, DEQF, DROPF)
```

CHECK WHETHER THE FLUID IN THE SHELL SIDE IS LIQUID OR VAPOR. С

С

IF(FLUIDF .EQ. 'LIQUID' .OR. FLUIDF .EQ. 'liquid') THEN GO TO 30 ELSEIF (FLUIDF .EQ. 'VAPOR' .OR. FLUIDF .EQ. 'vapor') THEN С CALCULATE THE HEAT-TRANSFER FILM COEFFICIENT IN THE VAPOR STATE. HTFF = HTFCV (CPF, GF, DEQF) ENDIF GO TO 90 IF (REF .LT. 2100.0) THEN 30 GO TO 50 ELSE HTFF = HTFCL (REF, PRF, KF, DEQF) ENDIF GO TO 90 50 HTFF = HTFCJ (REF, PRF, KF, DEQF, TL) CALCULATE THE HEAT-TRANSFER FILM COEFFICIENT CORRECTED C C FOR FOULING. 90 HIFF = HTFF/(1.0+HTFF*FOLF)С CALCULATE THE OVERALL HEAT-TRANSFER COEFFICIENT. С CALCULATE THE PARAMETER X, USED IN THE FIN EFFICIENCY X = FHT*(HIFF/(6.0*KFT*FTHICK))**0.5С CALCULATE THE FIN EFFICIENCY VAL1 = EXP(X) - EXP(-X)VAL2 = EXP(X) + EXP(-X) $FE = (1.0/X \star (VAL1/VAL2))$ CALCULATE THE EFFECTIVE SURFACE EFFICIENCY FOR FINS С ESEF = FE*(AF/AO)+(1.0-AF/AO)с CALCULATE THE CORRECTED FILM HEAT TRANSFER RATE $HIFD = HIFF \star ESEF$ CONVERT THE FIN EFFICIENCY AND EFFECTIVE SURFACE EFFICIENCY С С TO PERCENTAGES. FE = 100.0 * FE $ESEF = 100.0 \star ESEF$ с CALCULATE OVERALL HEAT TRANSFER COEFFICIENT. DELTW = (ODT/12.0) - (IDT/12.0)RATIO3 = (PI*IDT)/((PI*ODT)+ÁF) RATIO4 = (DELTW/KFT)+(1.0/(HIFF*RATIO3))+(1.0/HIFD) UO = 1.0/RATIO4

```
RETURN
     END
С
CCCC
     THIS PROGRAM CALCULATES THE LOG MEAN TEMPERATURE DIFFERENCE,
     LMTD (oF)
     SUBROUTINE LOGMTD (TH1, TH2, TC1, TC2, LMTD)
     REAL LMTD
     \begin{array}{rcl} \text{DELT1} &=& \text{TH1} &-& \text{TC2} \\ \text{DELT2} &=& \text{TH2} &-& \text{TC1} \end{array}
     LMTD = ABS((DELT1 - DELT2)/ALOG (DELT1/DELT2))
     RETURN
     END
     **********
C
C
     THIS SUBPROGRAM EVALUATES THE HEAT TRANSFER FILM COEFFICIENT
с
     IN THE VAPOR STATE OF A DOUBLE PIPE HEAT EXCHANGER.
      ċ
     FUNCTION HTFCV (CP1, G1, DEQ1)
     HTFCV = (0.0144*CP1*(G1**0.8))/(DEQ1**0.2)
     RETURN
     END
c
c
     THIS SUBPROGRAM EVALUATES THE HEAT TRANSFER FILM COEFFICIENT
c
c
     IN THE LIQUID STATE OF A DOUBLE PIPE HEAT EXCHANGER.
      **********
     FUNCTION HTFCL (RE1, PR1, K1, D1)
     REAL K1
     HTFCL = 0.023*(RE1**0.8)*(PR1**0.333)*(K1/D1)*1.0
     RETURN
     END
C
C
C
C
C
C
C
C
      THIS SUBPROGRAM EVALUATES THE HEAT TRANSFER FILM COEFFICIENT
     FOR REYNOLDS NUMBER LESS THAN 2100.0 OF A DOUBLE PIPE
     HEAT EXCHANGER
     FUNCTION HTFCJ (RE, PR, K1, D1, TL1)
     REAL K1
     HTFCJ = 1.86*(RE**0.333)*(PR**0.333)*(K1/D1)*(D1/TL1)**0.333)
     RETURN
     END
C
C
```

с с с с	CALCULATE THE FRICTION FACTOR IN THE SHELL SIDE OF THE DOUBLE PIDE HEAT EXCHANGER (Dennis Wright: Basic Programs for chemical engineers, Van Nostrand Reinhold company, NY 1986)
	SUBROUTINE FFACTOR (RE, FACTOR)
	IF (RE .GT. 2100) THEN
	FACTOR = 0.00033/(RE**0.25)
	ELSEIF (RE .LT. 2100) THEN
	FACTOR = 16.0/RE ENDIF RETURN END
с с с с	**************************************
	SUBROUTINE PDROP (FRICT, TLEN, NP, G, DEN, EQDIA, DROP)
	DROP = (4*FRICT*TLEN*NP*(G/3600.0)**2)/(64.4*DEN*EQDIA)
	RETURN END
C C C	**************************************
	SUBROUTINE RESULTS
	CHARACTER*30 FLUIDT, FLUIDS, FLUIDF, TUBE, FTTUBE, FTSHELL
	REAL IDTF, IDSF, IDFF, ODTF, ODSF, ODFF REAL IDT, ODT, KT REAL IDS, ODS, KS, KST REAL IDF, ODF, KF, KFT REAL LMTD
*	<pre>COMMON TUBE COMMON/DATA/FTTUBE, FTSHELL, TDEN, SDEN COMMON/DATA/FLUIDT, IDT, ODT, WT, CPT, VIST, KT, TT1, TT2, FOLT, TL, NTP COMMON/DATA2/FLUIDS, IDS, ODS, WS, CPS, VISS, KS, TS1, TS2, FOLS, KST, NSP COMMON/DATA3/FLUIDF, IDF, ODF, WF, CPF, VISF, KF, TF1, TF2, FOLF, FHT, FTHICK, NF, KFT, NFP COMMON/DATA5/FLUIDF, IDSF, IDFF, ODTF, ODSF, ODFF COMMON/DATA5/IDTF, IDSF, IDFF, ODTF, ODSF, ODFF COMMON/RES1/VELT, RET, PRT, FRICTT, DROPT, HTFT, HIFTU</pre>
*U	COMMON/RES2/VELS,RES,PRS,FRICTS,DROPS,HTFS,HIFS COMMON/RES3/VELF,REF,PRF,FRICTF,DROPF,HTFF,HIFF,X,FE,ESEF,HIFD,

WRITE (1, 120)

120 * *	FORMAT (/,18X,'DOUBLE PIPE HEAT EXCHANGER RATING',/,18X, 'USING BARE-TUBES/LONGITUDINAL FINNED TUBES'/,18X, 'FORCED CONVECTION WITH NO CHANGE OF PHASE')
130	WRITE (1, 130) Format (1H ,78(1H*))
140	WRITE (1, 140) FORMAT (45X,'TUBE SIDE', 10X, 'SHELL SIDE')
150	WRITE (1, 150) Format (78(1H-))
160	WRITE (1, 160) FTTUBE, FTSHELL Format (2X,'Fluid Name:', T50, A15, T65, A15)
	IF (TUBE .EQ. 'FINS' .OR. TUBE .EQ. 'fins') THEN
170	WRITE (1, 170) FLUIDT, FLUIDF FORMAT (2X,'TYPE OF PHASE FLOW:', T50, A15, T65, A15)
180	WRITE (1, 180) WT, WF FORMAT (2X,'FLUID FLOW RATE, lb/hr.:', T45, F10.0, T60, F10.0)
190 *	WRITE (1, 190) CPT, CPF FORMAT (2X,'SPECIFIC HEAT CAPACITY, Btu/Ib.oF:', T45, F8.3, T60, F8.3)
200	WRITE (1, 200) TDEN, SDEN FORMAT (2X,'FLUID DENSITY, 1b/ft^3.:', T45, F8.3, T60, F8.3)
210	WRITE (1, 210) VIST, VISF FORMAT (2X,'FLUID VISCOSITY, cP:', T45, F8.3, T60, F8.3)
220	WRITE (1, 220) KT, KF FORMAT (2X,'FLUID THERMAL COND., Btu/Ib.oF:', T45, F8.3, T60, F8.3)
230 *	WRITE (1, 230) TT1, TF1 FORMAT (2X,'FLUID INLET TEMPERATURE, oF:', T45, F8.3, T60, F8.3)
240 *	WRITE (1, 240) TT2, TF2 FORMAT (2X,'FLUID OUTLET TEMPERATURE, oF:', T45, F8.3, T60, F8.3)
250	WRITE (1, 250) IDT, IDF FORMAT (2X,'INSIDE DIAMETER, inch:', T45, F8.3, T60, F8.3)
260	WRITE (1, 260) ODT, ODF FORMAT (2X,'OUTSIDE DIAMETER, inch:', T45, F8.3, T60, F8.3)
270	WRITE (1, 270) FOLT, FOLF FORMAT (2X,'FOULING FACTOR:', T45, F8.3, T60, F8.3)
280 *	WRITE (1, 280) TL, NTP FORMAT (2X,'TUBE LENGTH, ft.:', T45, F8.3,/, 2X,'NUMBER OF TUBE PASSES:', T45, I2)

290	*	WRITE (1, 290) NF, FHT, FTHICK FORMAT (2X, 'NUMBER OF FINS:', T45, I2,/, 2X,'FIN HEIGHT, inch:', T45, F8.3,/, 2X,'FIN THICKNESS, inch:', T45, F8.3)
300	*	WRITE (1, 300) KFT FORMAT (2X,'THERMAL COND. TUBE MATERIAL, Btu/hr.ft.oF:', T45,F8.3)
310		WRITE (1, 310) VELT, VELF FORMAT (2X,'FLUID VELOCITY, ft/sec.:', T45, F8.3, T60, F8.3)
320		WRITE (1, 320) RET, REF FORMAT (2X,'REYNOLDS NUMBER:', T45, F9.0, T60, F9.0)
330		WRITE (1, 330) PRT, PRF FORMAT (2X,'PRANDTL NUMBER:', T45, F8.4, T60, F8.3)
340		<pre>WRITE (1, 340) FRICTT, FRICTF FORMAT (2X,'FRICTION FACTOR (ft^2/in^2):', T45, F8.6, T60, F8.6)</pre>
350		WRITE (1, 350) DROPT, DROPF FORMAT (2X,'PRESSURE DROP, psi.:', T45, F8.4, T60, F8.4)
360	*	<pre>WRITE (1, 360) X, FE, ESEF FORMAT (2X,'FIN MODULUS:', T45, F8.3,/, 2X,'FIN EFFICIENCY, %:', T45, F8.3,/, 2X,'EFFECTIVE SURFACE EFFICIENCY, %:', T45, F8.3)</pre>
370		WRITE (1, 370) LMTD FORMAT (2X,'LOG MEAN TEMP. DIFF., oF:', T45, F8.2)
380	*	<pre>WRITE (1, 380) HIFTU, HIFF FORMAT (2X,'HEAT TRANSFER FILM COEFF. CORRECTED',/,2X,</pre>
390	*	<pre>WRITE (1, 390) UO FORMAT (2X,'OVERALL HEAT TRANSFER COEFF., Btu/hr.ft^2.oF:',</pre>
400	*	WRITE (1, 400) Q, AREA FORMAT (2X,'HEAT LOAD ON UNIT, Btu/hr.:', T45, F12.1,/, 2X,'HEAT TRANSFER SURFACE ft^2.:', T45, F8.3)
		ELSEIF (TUBE .EQ. 'BARE-TUBE' .OR. TUBE .EQ. 'bare-tube') THEN
410		WRITE (1, 410) FLUIDT, FLUIDS FORMAT (2X,'TYPE OF PHASE FLOW:', T45, A15, T60, A15)
420		WRITE (1, 420) WT, WS FORMAT (2X,'FLUID FLOWRATE:, lb/hr.:', T45, F10.0, T60, F10.0)
430	*	WRITE (1, 430) CPT, CPS FORMAT (2X,'SPECIFIC HEAT CAPACITY, Btu/lb.oF:', T45, F8.3,T60, F8.3)
440		WRITE (1, 440) TDEN, SDEN FORMAT (2X,'FLUID DENSITY, 1b/ft^3:', T45, F8.3, T60, F8.3)
		WRITE (1, 450) VIST, VISS

450	FORMAT (2X,'FLUID VISCOSITY, cP:', T45, F8.3, T60, F8.3)
460 *	WRITE (1, 460) KT, KS FORMAT (2X,'FLUID THERMAL COND., Btu/Ib.oF:', T45, F8.3, T60, F8.3)
470	WRITE (1, 470) TT1, TS1 FORMAT (2X,'FLUID INLET TEMPERATURE, oF:',T45, F8.3, T60, F8.3)
480	WRITE (1, 480) TT2, TS2 FORMAT (2X,'FLUID OUTLET TEMPERATURE, oF:',T40, F8.3,T60,F8.3)
490	WRITE (1, 490) IDT, IDS FORMAT (2X,'INSIDE DIAMETER, inch:',T45, F8.3, T60, F8.3)
500	WRITE (1, 500) ODT, ODS FORMAT (2X,'OUTSIDE DIAMETER, inch:', T45, F8.3, T60, F8.3)
510	WRITE (1, 510) FOLT, FOLS FORMAT (2X,'FOULING FACTOR:', T45, F8.3, T60, F8.3)
520 *	WRITE (1, 520) KFT FORMAT (2X,'THERMAL COND. TUBE MATERIAL, Btu/hr.ft.oF:', T45, F8.3)
530	WRITE (1, 530) VELT, VELS FORMAT (2X,'FLUID VELOCITY, ft/sec.:', T45, F8.3, T60, F8.3)
540	WRITE (1, 540) RET, RES FORMAT (2X,'REYNOLDS NUMBER:', T45, F9.0, T60, F9.0)
550	WRITE (1, 550) PRT, PRS FORMAT (2X,'PRANDTL NUMBER:', T45, F8.3, T60, F8.3)
560	WRITE (1, 560) FRICTT, FRICTS FORMAT (2X,' FRICTION FACTOR (ft^2/in^2):', T45, F8.5,T60,F8.5)
570	WRITE (1, 570) DROPT, DROPS FORMAT (2X,'PRESSURE DROP, psi:', T45, F8.4, T60, F8.4)
580	WRITE (1, 580) LMTD FORMAT (2X,'LOG MEAN TEMP. DIFF., oF:', T45, F8.2)
590 *	<pre>WRITE (1, 590) HIFTU, HIFS FORMAT (2X,'HEAT TRANSFER FILM COEFF. CORRECTED',/,2X,</pre>
600 *	<pre>WRITE (1, 600) UO FORMAT (2X,'OVERALL HEAT TRANSFER FILM COEFF., Btu/hr.ft^2.oF:',</pre>
610 *	WRITE (1, 610) Q, AREA FORMAT (2%,'HEAT LOAD ON UNIT , Btu/hr.:', T45, F12.1,/, 2%,'HEAT TRANSFER SURFACE, ft^2.:', T45, F8.3)
	ENDIF
620	WRITE (1, 620) FORMAT (1H ,78(1H-))
с	FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
	WRITE (1, *) CHAR(12)
	RETURN END

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PROGRAM PROG84

~	
С	***************************************
с	* THE SIZING OF AIR-COOLED HEAT EXCHANGERS.
с	* THE PROGRAM PERFORMS PRELIMINARY AIR-COOLER DESIGN CALCULATIONS
С	* WITH A RANGE OF INPUT DATA.
с	*
с	* THE PROGRAM CALCULATES:
c	* 1. THE NUMBER OF TUBE ROWS (TUBE BUNDLE DEPTH).
č	* 2. OUTLET AIR TEMPERATURE.
č	* 3. LOG MEAN TEMPERATURE DIFFERENCE.
c	
	to follow bind ford bonting inchi (bibbb on f filon or fobing
с	* ON 2 3/8 inch TRIANGULAR SPACING.).
С	* 5. COOLER FACE AREA (plan area).
с	* 6. AIR VELOCITY ACROSS TUBES.
с	* 7. TOTAL AIR FLOW.
с	* 8. FAN HORSEPOWER (bhp = brake horsepower).
с	* 9. TOTAL BUNDLE WEIGHT.
С	* 10. TUBE BUNDLE WIDTH FOR A GIVEN TUBE LENGTH.
С	
С	* THIS PROGRAM IS BASED ON CORRELATIONS PRESENTED IN ARTICLES BY
С	* E.C. Smith AND R. Brown. THE CORRELATIONS PRESENTED BY GRAPHS
С	* AND TABLES IN THE ORIGINAL ARTICLES AND FITTED TO A SERIES OF
с	* EQUATION BY W.W. Blackwell ARE USED IN THIS PROGRAM.
Ċ	*************
c	*
č	* R = NUMBER OF TUBE ROWS.
č	* TPI = INLET PROCESS FLUID TEMPERATURE, oF.
č	 * TAI = INLET AIR TEMPERATURE, OF. * TAI = INLET AIR TEMPERATURE, OF.
c	
с	
с	* $FV = FACE VELOCITY OF AIR, ft^2.$
с	* TAO1 = ESTIMATED AIR OUTLET TEMPERATURE, oF.
С	* U = OVERALL HEAT-TRANSFER COEFFICIENT
с	(BASED ON BARE-TUBE OD) Btu/hr.ft^20F.
с	* TPO = OUTLET PROCESS FLUID TEMPERATURE, oF.
С	* LMTD = EFFECTIVE LOG MEAN TEMPERATURE DIFFERENCE, oF.
С	* $Q = HEAT LOAD, Btu/hr.$
С	* TAO = CALCULATED OUTLET AIR TEMPERATURE, oF.
С	* F = AIR FLOW OVER TUBES, std ft^3/min.
с	* BHP = FAN HORSEPOWER, brake horsepower.
с	* WT = AIR-COOLER WEIGHT, 1b.
с	* W = TUBE BUNDLE, WIDTH, ft.
с	* TL == TUBE LENGTH, ft.
С	* C1,C2,C3,C4,C5,C6,C7,C8,C9,C10 = CONSTANTS
с	********
	REAL LMTD
	-
	OPEN (UNIT=3, FILE='DATA84.DAT', STATUS='OLD', ERR=18)
	OPEN (UNIT =1, FILE='PRN')
с	CONSTANTS USED IN THE PROGRAM
~	
	C1 = 3.1679

C1 = 3.1679C2 = 3.7948

```
C3 = 1.2557
          C4 = 1.0031
          C5 = 720.8542
          C6 = 0.9530
          C7 = 7.4212
          C8 = 12.5342
          C9 = 36.4
          C10 = 9.35
С
с
с
с
с
          READ THE INLET PROCESS FLUID TEMPERATURE, OF: TPI
          READ THE OUTLET PROCESS FLUID TEMPERATURE, OF: TPO
          READ THE INLET AIR TEMPERATURE, OF: TAI
READ THE OVERALL HEAT-TRANSFER COEFFICIENT, Btu/hr.ft^2.oF: U
с
          READ THE HEAT LOAD, Btu/hr.: Q
č
          READ THE TUBE LENGTH, ft.: TL
         READ (3, *, ERR=19) TPI, TPO, TAI
READ (3, *, ERR=19) U, Q, TL
GO TO 20
 18
          WRITE (*, 100)
 100
          FORMAT (6X, 'DATA FILE DOES NOT EXIST')
          GO TO 999
          WRITE (*, 110)
FORMAT (6X, 'ERROR MESSAGE IN THE DATA VALUE')
 19
 110
          GO TO 999
 20
          WRITE (1, 120)
 120
          FORMAT (///,20X,'AN AIR-COOLED HEAT EXCHANGER DESIGN',/,78(1H*))
          WRITE (1, 130) TPI, TPO, TAI
         FORMAT (6X,'INLET PROCESS FLUID TEMPERATURE, oF:',T60, F8.1,
/,6X,'OUTLET PROCESS FLUID TEMPERATURE, oF:',T60,F8.1,
/,6X,'INLET AIR TEMPERATURE, oF:',T60,F8.1)
 130
      *
         WRITE (1, 140) U, Q, TL
FORMAT (6X,'OVERALL HEAT-TRANSFER COEFFICEINT, Btu/hr.ft^2.oF:',
 140
                    T60, F8.1,/,6X,'HEAT LOAD, Btu/hr.:',T60, F12.0,/,6X,
'TUBE LENGTH, ft.:', T60, F8.1)
      ٠
      *
          CALCULATE THE NUMBER OF TUBE ROWS
С
          R = C1 + C2*(ALOG((TPI-TAI)/U))
          CALCULATE THE RATIO OF BARE-TUBE SURFACE AREA BASED ON THE
С
ċ
          TUBE OD TO THE FACE AREA OF BUNDLE , RATIO1=BTSA/FA
          RATIO1 = C3*(R**C4)
с
          CALCULATE THE FACE VELOCITY OF AIR, ft/min.
```

FV = C5*(C6**R)

.

WRITE (1, 150) R, RATIO1, FV WRITE (1, 190, R, KAILOT, T FORMAT (6X, 'NUMBER OF TUBE ROWS:', T60, F6.1, /, 6X, 'RATIO OF BARE-TUBE SURFACE TO FACE AREA OF BUNDLE', T60, F6.1, 'RATIO OF BARE-TUBE SURFACE TO FACE AREA OF BUNDLE', T60, F6.1, 150 /, 6X, 'FACE VELOCITY OF AIR, ft./min.:', T60, F8.3) С CALCULATE THE ESTIMATED AIR OUTLET TEMPERATURE, OF: TAO1 TAO1 = (0.005 * U * ((TPO + TPI) / 2.0 - TAI)) + TAIWRITE (1, 160) TAO1 160 FORMAT (6X, 'ESTIMATED AIR OUTLET TEMPERATURE, oF:', T60, F8.3) С CALCULATE THE LOG MEAN TEMPERATURE DIFFERENCE, oF: LMTD. 25 VAL1 = (TPO-TAI) - (TPI-TAO1)VAL2 = ALOG((TPO-TAI)/(TPI-TAO1)) LMTD = VAL1/VAL2 С CALCULATE BARE-TUBE SURFACE AREA BASED ON TUBE, OD: ft^2: BTSA BTSA = Q/(U*LMTD)С CALCULATE THE FACE AREA OF BUNDLE, ft^2: FA FA = BTSA/RATIO1 WRITE (1, 170) LMTD, BTSA, FA FORMAT (6X,'EFFECTIVE LOG MEAN TEMPERATURE DIFFERENCE, oF:',T60, 170 F8.3, /, 6X, 'BARE-TUBE SURFACE AREA BASED ON TUBE OD, ft^2:', T60, F8.1, /, 6X, 'FACE AREA OF BUNDLE, ft^2:', T60, F8.1) С CALCULATED OUTLET AIR TEMPERATURE, OF: TAO TAO = (Q/(FA*FV*1.08)) + TAIWRITE (1, 180) TAO 180 FORMAT (6X, 'CALCULATED OUTLET AIR TEMPERATURE, oF:', T60, F8.3) С A NECESSARY CONDITION FOR SIZING AN AIR-COOLED EXCHANGER. с CHECK WHETHER THE DIFFERENCE BETWEEN THE CALCULATED AIR OUTLET TEMPERATURE AND THE ESTIMATED AIR OUTLET TEMPERATURE IS GREATER С с THAN/OR EQUAL TO 0.5 (i.e. (TAO-TAO1) .GE. 0.5)) с IF THE DIFFERENCE IS GREATER THAN OR EQUAL TO 0.5, THEN SET THE ESTIMATED AIR OUTLET TEMPERATURE EQUAL TO THE CALCULATED AIR С OUTLET TEMPERATURE AND REPEAT THE CALCULATION FROM THE LOG MEAN С C TEMPERATURE DIFFERENCE. IF ((TAO-TAO1) .GE. 0.5) THEN WRITE (1, 190) FORMAT (6X,'THE DIFFERENCE BETWEEN THE CALCULATED AIR', / 'OUTLET TEMPERATURE AND THE ESTIMATED AIR OUTLET', /, 6X, 'TEMPERATURE IS GREATER THAN 0.5',T60,'TAO-TAO1 >0.5') 190 /, 6X, * TAO1 = TAOGO TO 25 ELSE

ELSE ENDIF

```
200
     *
     ÷
с
         CALCULATE AIR FLOW OVER TUBES, std ft^3/min.: F
         F = FA * FV
С
         CALCULATE FAN HORSEPOWER: BHP
         BHP = BTSA/(C7+(C8*R))
         CALCULATE AIR-COOLER WEIGHT, 1b: WT.
С
         WT = FA*(C9+(C10*R))
        CALCULATE TUBE BUNDLE WIDTH, ft: W
С
         W = FA/TL
С
        WRITE (1, 210) F, BHP
FORMAT (6X,'AIR FLOW OVER TUBES, std.ft^3/min.:', T60, F9.0,
/, 6X, 'FAN HORSEPOWER, bhp:', T60, F8.1)
 210
     *
        WRITE (1, 220) WT, W
FORMAT (6X,'AIR-COOLER WEIGHT, 1b:', T60, F9.1, /, 6X,
'TUBE BUNDLE WIDTH, ft.:',T60,F8.1)
 220
     ٠
         WRITE (1, 230)
 230
         FORMAT (1H ,78(1H-))
с
         FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
         WRITE (1, *) CHAR(12)
         CLOSE (3, STATUS='KEEP')
         CLOSE (1)
  999
         STOP
         END
```

PROGRAM PROG85

с с с с с с с с	THE PROG FOR ANY MEDIUM A	CER REQUIREMENTS FOR PIPELINES. TRAM CALCULATES THE HEAT LOSS AND TRACING REQUIREMENTS GIVEN PIPELINE USING HOT OIL OR SOME OTHER HOT FLUID IS THE HEAT SOURCE. IT ALSO CALCULATES THE HEAT LOSS INSULATED PIPELINE WITHOUT A HEAT TRACER.
с с с с с с с с	1. SURF 2. TOTA 3. TOTA 4. FLOW 5. TOTA	GRAM WILL CALCULATE: ACE TEMPERATURE OF INSULATED TRACED PIPE. L HEAT TRANSFERRED PER 100 ft. OF PIPE. L HEAT TRANSFERRED FOR ENTIRE PIPELINE. RATE FOR HOT MEDIA. L NUMBER OF HEAT TRACERS REQUIRED WITH AND WITHOUT ISFER CEMENT.
c c	HEAT TRA RESISTAN	NSFERRED THROUGH AN INSULATED PIPE ENCOUNTERS FOUR ICES:
0000	2. HEAT 3. HEAT	RESISTANCE ON THE INSIDE WALL OF THE PIPE. RESISTANCE THROUGH THE PIPE WALL. RESISTANCE THROUGH THE INSULATION. FILM RESISTANCE ON THE OUTSIDE OF THE INSULATION.
0000	VERY SMA	THE FIRST TWO RESISTANCES, 1. AND 2. ARE NORMALLY ALL AND ARE THEREFORE NEGLIGIBLE IN THIS PROGRAM.
с с с	TMI	 AVERAGE TEMPERATURE OF HOT MEDIUM, oF. HOT-MEDIUM INLET TEMPERATURE, oF. HOT-MEDIUM OUTLET TEMPERATURE, oF.
C C	T AVG DI	= AVERAGE TEMPERATURE OF PIPE AND TRACER, oF. = INSIDE DIAMETER OF INSULATION, in.
C C C	TAL	 OUTSIDE DIAMETER OF PIPE, in. ALLOWANCE FOR TRACER DIAMETER, in. OUTSIDE DIAMETER OF INSULATION, in.
c c	TK O	= INSULATION THICKNESS, in. = HEAT LOSS PER FOOT OF PIPE, Btu/hr.ft.
c c	KINS TS	<pre>= THERMAL CONDUCTIVITY OF INSULATION, Btu/hr.ft^2.(oF/ft). = OUTSIDE SURFACE TEMPERATURE OF INSULATION, oF.</pre>
с с с		FILM HEAT-TRANSFER COEFFICIENT TO AIR CORRECTED FOR THE WIND, Btu/hr.ft^2.oF. AIR TEMPERATURE, oF.
c c	HCOMB	= HC+HR: COMBINED CONVECTIVE AND RADIATIVE HEAT TRANSFER CO-EFFICIENT, Btu/hr.ft^2.oF.
c c		= WIND FACTOR.
c		<pre>= TOTAL HEAT LOSS FROM PIPELINE, Btu/hr. = FLOW RATE OF HOT MEDIUM, lb/hr.</pre>
č		= TOTAL PIPELINE LENGTH, ft.
с		= SPECIFIC HEAT OF HOT MEDIUM, Btu/lb.oF.
С	TWOC	= NUMBER OF TRACERS REQUIRED WITHOUT HEAT-TRANSFER CEMENT.
c c		= THERMAL CONDUCTANCE, TRACER TO PIPE WITHOUT CEMENT, Btu/hr.oF(ft of pipe).
с		= NUMBER OF TRACERS REQUIRED WITH HEAT TRANSFER CEMENT.
С	TCB	= THERMAL CONDUCTANCE, TRACER TO PIPE
c c	TР	WITH CEMENT, Btu/hr.oF(ft of pipe). = TEMPERATURE IN THE PIPE, oF.
c	-	- TEMPERATURE IN THE FIFE, OF.
-		

	REAL OD, KINS
	OPEN (UNIT=3, FILE ='DATA85.DAT', STATUS='OLD', ERR=18) OPEN (UNIT=1, FILE ='PRN')
	PI = 3.1415927
000000000000000000	READ THE TOTAL LENGTH OF PIPE, ft.: TOTL READ THE TEMPERATURE IN THE PIPE, oF: TP READ THE HOT-MEDIUM INLET TEMPERATURE, oF: TMI READ THE HOT-MEDIUM OUTLET TEMPERATURE, oF: TMO READ THE AIR TEMPERATURE, oF: TAIR READ THE AIR TEMPERATURE, oF: TAIR READ THE ALLOWANCE FOR THE TRACER DIAMETER, inch.: TAL READ THE ALLOWANCE FOR THE TRACER DIAMETER, inch.: TAL READ THE INSULATION THICKNESS, inch.: TK READ THE INSULATION THICKNESS, inch.: TK READ THE SPECIFIC HEAT CAPACITY OF THE HOT MEDIUM, Btu/h.ft^2(oF/ft): KINS READ THE FILM HEAT TRANSFER COEFFICIENT TO AIR CORRECTED FOR THE WIND, Btu/hr.ft ² .oF: HA READ THE THE THERMAL CONDUCTANCE, TRACER TO PIPE WITHOUT CEMENT, Btu/hr.oF (ft. of pipe).: TCA READ THE THERMAL CONDUCTANCE, TRACER TO PIPE WITH CEMENT, Btu/hr.oF (ft. of pipe).: TCB
	READ (3, *, ERR=19) TOTL, TP, TMI READ (3, *, ERR=19) TMO, TAIR, OD READ (3, *, ERR=19) TAL, TK, KINS READ (3, *, ERR=19) CP, HA, TCA, TCB
	GO TO 20
18 100	WRITE (*, 100) Format (6X, 'data file does not exist') Go to 999
19 110	WRITE (*, 110) Format (6x, 'error message in the data value') Go to 999
20 120 *	<pre>WRITE (1, 120) FORMAT (///, 10X, 'HEAT TRACER REQUIREMENTS FOR PIPELINES AND ', /, 10X, 'HEAT LOSS FROM AN INSULATED PIPELINE WITH TRACING', /, 78(1H*))</pre>
130 * *	WRITE (1, 130) TOTL, TP, TMI FORMAT (6X, 'TOTAL PIPELINE LENGTH, ft.:',T60,F9.1,/, 6X, 'TEMPERATURE IN PIPE, oF:',T60,F9.1,/, 6X, 'HOT-MEDIUM INLET TEMPERATURE, oF:',T60,F9.1)
140 * *	<pre>WRITE (1, 140) TMO, TAIR, OD FORMAT (6X, 'HOT MEDIUM OUTLET TEMPERATURE, oF:',T60, F9.1,/, 6X, 'AIR TEMPERATURE, oF:',T60,F9.1,/, 6X, 'OUTSIDE DIAMETER OF PIPE, in.:',T60,F9.3)</pre>
150 * *	<pre>WRITE (1, 150) TAL, TK, KINS FORMAT (6X, 'ALLOWANCE FOR TRACER DIAMETER, in.:', T60, F9.2,/, 6X, 'INSULATION THICKNESS, in.:', T60, F9.2, /, 6X, 'THERMAL CONDUCTIVITY OF INSULATION, Btu/hr.ft^2(oF/ft):', T60,</pre>

* F9.3)

160 * * * *	/,6X,'THERMAL CONDUCTANCE TRACER TO PIPE', /,6X,'WITHOUT CEMENT, Btu/hr.oF.ft of pipe:',T60,F9.3
с	CALCULATE THE AVERAGE TEMPERATURE OF HOT MEDIUM, OF
	CONTRL = 0.0
30	TMEDAV = 0.5*(TMI+TMO) TA = 0.5*(TMEDAV+TP) DI = OD+TAL DO = DI+2.0*TK HA1 = HA
40	X = (ALOG(DO/DI)*HA1*DO)/(24.0*KINS) TS = (TA+X*TAIR)/(1.0+X)
170 *	WRITE (1, 170) TS FORMAT (6X, 'OUTSIDE SURFACE TEMPERATURE OF INSULATION, oF:', T60,F9.3)
C C	CALCULATE THE COMBINED CONVECTIVE AND RADIATIVE HEAT TRANSFER COEFFICIENT.
	HCOMB = 564.0/((DO**0.19)*(273.0-(TS-TAIR)))
180 *	WRITE (1, 180) HCOMB FORMAT (6X,'THE COMBINED CONVECTIVE AND RADIATIVE', /, 6X, 'HEAT TRANSFER COEFFICIENT, Btu/h.oF.ft^2:', T60, F9.3)
с	CONSTANTS FOR THE WIND FACTOR
	C0 = 2.814 C1 = -0.00038857 C2 = -0.0000012857
	WF = CO+(C1*(TS-TAIR))+(C2*((TS-TAIR)**2))
c c	CALCULATED FILM HEAT-TRANSFER COEFFICIENT TO AIR CORRECTED FOR THE WIND, HA2: Btu/hr.ft^2.oF
	$HA2 = HCOMB \star WF$
	IF ((HA2-HA1) .GE. 0.01) THEN HA1 = HA2 GO TO 40 ELSE
с	CALCULATE THE HEAT FLOW THROUGH THE INSULATION Btu/hr.(lin ft).
	QINS = (2.0*PI*KINS*(TA-TS))/(ALOG(DO/DI))

с CALCULATE THE HEAT FLOW BY RADIATION AND CONVECTION TO AIR QRC = (HA1*PI*DO*(TS-TAIR))/12.0ENDIF WRITE (1, 190) WF, HA2 FORMAT (6X,'WIND FACTOR:',T60, F9.3, /, 6X, 'HEAT TRANSFER COEFFICIENT TO AIR', /, 6X, 'CORRECTED FOR THE WIND, Btu/hr.ft^2.oF:', T60, F9.3) 190 CALCULATE THE TOTAL HEAT LOST FROM PIPELINE ,Btu/hr. С QT = QINS*TOTLWRITE (1, 200) QINS, QT FORMAT (6X,'HEAT LOSS PER FOOT OF PIPE, Btu/hr.ft:', T60,F9.3,/, 6X,'TOTAL HEAT LOSS FROM PIPELINE, Btu/hr.:', T60, F9.3) 200 IF (CONTRL .EQ. 1.0) THEN GO TO 999 ELSE CALCULATE THE FLOW RATE OF HOT MEDIUM, 1b/h С W = QT/(CP*(TMI-TMO))WRITE (1, 210) W 210 FORMAT (6X, 'FLOW RATE OF HOT MEDIUM, 1b/hr.:', T60, F9.3) CALCULATE THE NUMBER OF TRACERS REQUIRED WITHOUT С HEAT-TRANSFER CEMENT. C TWOC = QINS/(TCA*(TMEDAV-TP)) CALCULATE THE NUMBER OF TRACERS REQUIRED WITH С HEAT-TRANSFER CEMENT. с TWC = QINS/(TCB*(TMEDAV-TP)) ENDIF WRITE (1, 220) TWOC, TWC FORMAT(6X,'THE NUMBER OF TRACERS WITHOUT HEAT TRANSFER CEMENT:', 220 'THE NUMBER OF TRACERS WITH HEAT TRANSFER CEMENT:', T60, F9.2) * WRITE (1, 230) 230 FORMAT (1H ,78(1H-)) С ******* С CALCULATION OF HEAT LOSS FROM AN INSULATED PIPE WITHOUT С TRACING, WHERE TMI=TMO=TP, AND TA=0.0 WRITE (1, 240) FORMAT (//, 6X, 'HEAT LOSS FROM AN INSULATED PIPE', \) 240 WRITE (1, 250) FORMAT (' WITHOUT TRACING', /1H ,78(1H*)) 250

C TO CALCULATE THE HEAT LOSS FROM AN INSULATED PIPE, CONTRL=1.0 TMI = TP TMO = TP TAL = 0.0 CONTRL = CONTRL + 1.0 GO TO 30 CLOSE (3, STATUS='KEEP') CLOSE (1) 999 STOP END

CHAPTER 9

Engineering Economics

INTRODUCTION

Project evaluation enables the technical and economic feasibility of a chemical process to be assessed using preliminary process design and economic evaluations. Once a process flowsheet is available, these evaluations can be classified into several steps: material balance calculations, equipment sizing, equipment cost determination, utilities requirements, investment cost estimation, sales volume forecasting, manufacturing cost estimation, and finally profitability and sensitivity analysis.

The results of an economic evaluation are reviewed together with other relevant aspects, such as, competition and likely product life in arriving at project investment decisions. The decisions are essential in order to both plan and allocate the long-term use of available resources. The objectives of an economic appraisal are:

- To ensure that the expected future benefits justify the expenditure of resources.
- To choose the best project from among alternatives to achieve future benefits.
- To use all available resources.

These objectives ensure that the right project is undertaken and is likely to attain the desired profitability.

In the chemical process industries, an investment project may arise from any of a range of activities. It may be a minor modification to an existing plant, a major plant expansion or revamping, a completely new plant (on an existing or greenfield site) or the development of an entirely new process or product. For a major or a new plant, economic assessment with increasing degrees of accuracy may be carried out at different stages as the plant progresses. This may be based on an initial research and development (R&D), through various stages of the project (e.g., pilot plants, material evaluation, preliminary plant design), which leads to the decision whether or not to proceed with investment in a full scale plant for the process. Table 9-1 lists examples of the use of engineering economics. The approach used for economic evaluation depends on the quality of the information, that is, the stage in the project at which it is undertaken.

Table 9-1Use of Chemical Engineering Economics

Production or Plant Technical Services

- A. Plant equipment continuously needs repair, replacement, or modernization. The responsible engineer should know roughly what the comparative performance, costs, and payout periods are, even if there is a plant engineering group to do that type of analysis, or a firm price quotation will be obtained later.
- B. Plant changes, such as those initiated by the ever increasing costs of energy and environmental requirements, necessitate that many energy saving, pollution, and hazardous waste control possibilities must be considered. To make intelligent recommendations, the responsible engineer should personally conduct design and cost estimates, pay-back, and economic calculations on the alternatives before making even preliminary recommendations.
- C. Competitors' processing methods, as well as R&D, sales, or management suggested changes must be continuously examined. Supervisors or other groups may be responsible, but the staff engineer can help a great deal by making preliminary cost estimates and economic analyses of the changes to guide his or her own thinking (and hopefully the group's position).
- D. All engineers should have a feel for their company's business, products, and economics. This requires occasional economic reading, and a basic understanding of the company's annual reports and general industry economic news.

Research and Development

A. In the process of being creative, one thinks of many novel ideas. During the analytical phase of creative thinking many of the ideas will require a quick cost estimation and economic analysis to provide a better idea of their merit. Even though supervisors or others may be assigned to do this work, the chances of conceiving good ideas and having them accepted increases immensely if some economic screening can be done by the originator.

- B. While conducting R&D studies, there are always many stumbling points or alternative directions that may be taken to solve problems. Often brief cost estimates and economic analyses will help the engineer in deciding which are the most promising directions to pursue.
- C. After an early or intermediate stage of an R&D program has been successful, new funding requests usually are required to continue the study. These requests can always benefit from having potential preliminary economic analyses. Later, in the final stages of a successful project, the engineer may be part of a team assigned to provide a more definitive preliminary economic projection and analysis.
- D. In dealing with production, sales, or management personnel, one can usually gain more respect, and be considered more practical and less "theoretical," by having a reasonable knowledge of the costs and economics of the projects under study, and general industry economics.

Sales

- A. A general knowledge of company costs, profits, and competition are very helpful for more effective salesmanship.
- B. Salesmen often recommend new products, improvements, or pricing ideas to their management. A cost and economic estimate for these ideas should be helpful in the proposal report.
- C. Salesmen sometimes perform market surveys. Again, a general economic knowledge of the industries and companies surveyed may be essential, and is always useful.
- D. Salesmen may move into management, where economic knowledge is a major part of the job.

Engineering

A. Because of the extreme specialization of most engineering companies and many company engineering departments, cost estimating and economics may not be directly required in many engineering company or department jobs. However, other jobs will deal exclusively with cost estimating and economic analysis, and all will benefit from a good, fluent knowledge of the basic economic procedures. Engineering departments or companies usually have well developed in-house methods and data that must be used, but the basics are still applicable.

General

A. All chemical engineers are assumed to know the rudiments of cost estimating, economic evaluation, and the economics of their industry. A high percentage will find this knowledge useful or necessary throughout their careers.

(table continued on next page)

Table 9-1 (continued)

B. All work situations are more or less competitive, and one means of maintaining the highest advancement potential with most jobs is to convince superiors of your knowledge and interest in management, business, and economics. Associated with this is the demonstration of ability, an interest in accepting responsibility, and ability to communicate. Many companies promote people whom they think can "manage," such as those with MBA (master of business administration) degrees or with perceived equivalent capabilities, ahead of people with a better performance record. Basically, a confidence that you can learn managerial skills as you need them and a knowledge of economics should make most chemical engineers equal or preferred candidates for advancement.

Source: Garrett [10], Courtesy of Chapman and Hall

Project Evaluation

Investment decisions are often based upon several criteria, such as annual return on investment (ROI), payback period (PBP), net present value (NPV), the average rate of return (ARR), present value ratio (PVR), or the internal rate of return (IRR). Discounted cash flow rate on return (DCFRR) is another popular means of evaluating the economic viability of a proposed project. Horwitz [1] recommended the DCFRR as the best means to determine the return on investment, because it accounts for the time value of money. The internal rate of return as an investment criterion gives the possibility that given cash flows may result in more than one internal rate of return. Cannaday *et al.* [2] developed a method for determining the relevance of an internal rate of return. They inferred that an internal rate of return is relevant, if its derivative with respect to each of the cash flows is positive.

Powell [3] reviewed the basics of various discounted cash flow techniques for project evaluation. Discounting is a method that accounts for the time value of money to provide either for the capital or to convert the cash flows to a common point in time so that they are summed. Ward [4] proposed a new concept known as the net return rate (NRR) that provides a better indication of a project's profitability. Techniques and criteria for economic evaluation of projects are widely available in the literature and texts [5,6,7,8,9,10]. A summary of the conventional decision criteria is given here.

Cash Flows

Return On Investment (ROI)

In engineering economic evaluation, rate of return on investment is the percentage ratio of average yearly profit (net cash flow) over the productive life of the project, divided by the total initial investment. This is calculated after income taxes have been deducted from the gross or pre-tax income. The remainder or net income may be used either for paying dividends, reinvestment, or can be spent for other means. ROI is defined by

$$ROI = \frac{Annual return}{Investment} \times 100$$
(9-1)

The annual return may be the gross income, net pre-tax income, net after-tax income, cash flow, or profit. These may be calculated for one particular year or as an average over the project life. Investment may be the original total investment, depreciated book-value investment, lifetime average investment, fixed capital investment, or equity investment. The investment includes working capital and sometimes capitalized expenses such as interest on capital during construction.

Payback Period (PBP)

Payback period is widely used when long-term cash flows are difficult to forecast, because no information is required beyond the break-even point. It may be used for preliminary evaluation or as a project screening device for high risk projects in times of uncertainty. Payback period is usually measured as the time from the start of production to recovery of the capital investment. The payback period is the time taken for the cumulative net cash flow from start-up of the plant to equal the depreciable fixed capital investment ($C_{FC} - S$). It is the value of t that satisfies the equation

$$\sum_{t=0}^{t=(PBP)} C_{CF} = (C_{FC} - S)$$
(9-2)

where C_{CF} = net annual cash flow C_{FC} = fixed capital cost S = salvage value

Figure 9-1 shows the cumulative cash flow diagram for a project. The PBP is the time that elapses from the start of the project, A, to the

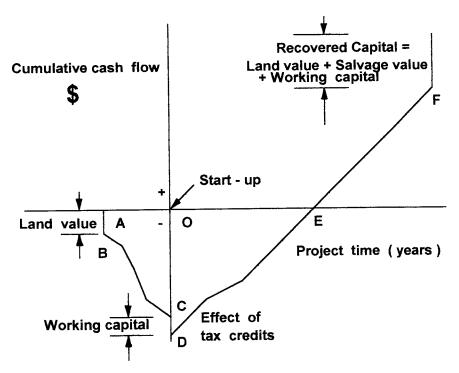


Figure 9-1. Cumulative cash flow diagram.

break-even point, E, where the rising part of the curve passes the zero cash position line. The PBP thus measures the time required for the cumulative project investment and other expenditure to be balanced by the cumulative income.

Present Worth (Or Present Value)

In an economic evaluation of a project, it is often necessary to evaluate the present value of funds that will be received at some definite time in the future. The present value (PV) of a future amount can be considered as the present principal at a given rate and compounded to give the actual amount received at a future date. The relationship between the indicated future amount and the present value is determined by a discount factor. Discounting evaluates each year's flow on an equal basis. It does this by means of the discount, or present value factor, and the reciprocal of the compound interest factor $(1 + i)^n$ with

i = interest rate

n = the year in which the interest is compounded

The discount factor
$$= \frac{1}{(1+i)^n}$$
 (9-3)

If C_n represents the amount available after n interest periods, P is the initial principal and the discreet compound interest rate is i. The present value, PV, can be expressed as:

$$PV = P = \frac{C_n}{(1+i)^n}$$
 (9-4)

Net Present Value (NPV)

The net present value of a project is the cumulative sum of the discounted cash flows including the investment. The NPV corresponds to the total discounted net return, above and beyond the cost of capital and the recovery of the investment. The NPV represents a discounted return or profit, but is not a measure of the profitability. Each cash flow is evaluated by computing its present value. This is done by taking a cash flow of year n and multiplying it by the discount factor for the nth year.

Present value of
$$p_n = C_n \left[\frac{1}{(1+i)^n} \right]$$
 (9-5)

The present value, p, at year 0 of a cash flow, C_t , in year t at an annual discount rate of i is

$$p = \frac{C_t}{\left(1+i\right)^t} \tag{9-6}$$

For a complete project, the earlier cash flows are usually negative and the later ones positive. The net present value, NPV, is the sum of the individual present values of the yearly cash flows. This is expressed as:

NPV =
$$C_0 + \frac{C_1}{(1+i)} + \frac{C_2}{(1+i)^2} + \dots + \frac{C_n}{(1+i)^n}$$
 (9-7)

Equation 9-7 can be expressed as:

NPV =
$$\sum_{t=0}^{t=n} \frac{C_t}{(1+t)^t}$$
 (9-8)

The life of the project n years must be specified together with the estimated cash flows in each year up to n. NPV = net present value C_0 = Initial investment C_n = cash flow n = year n i = interest rate of return (ROI/100)

If a project includes a series of identical yearly cash flows, such as the net inflow for several years, their present values can be obtained in one calculation instead of singly. For yearly cash flows, C, from year m to year n, then the sum of their present values is:

$$P = \frac{C}{i(1+i)^{m-1}} \left[1 - \frac{1}{(1+i)^{n-m+1}} \right]$$
(9-9)

Figure 9-2 shows the cumulative net present values stages in a project. Tables 9-2 through 9-5 are the discount factors for computing the net present value.

Assuming that the investment is made in year 0 ($C_0 = I$), and the cash flows over the project life are constant, then Equation 9-7 is simplified to give

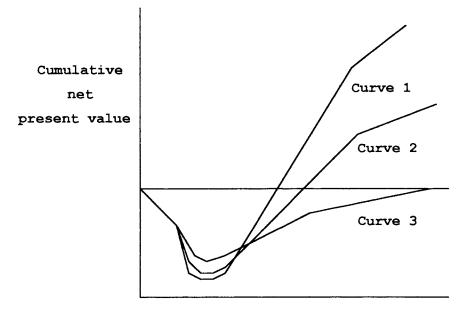
$$NPV = \left[C\sum_{n=1}^{n} \left(\frac{1}{1+i}\right)^{n}\right] - I$$
(9-10)

The term 1/(1+i) is a geometric progression whose sum can be expressed as the single term

$$\sum_{n=1}^{n} \frac{1}{(1+i)^{n}} = \frac{(1+i)^{n} - 1}{i(1+i)^{n}}$$
(9-11)

where Equation 9-11 becomes the present value of an annuity. Equation 9-7 can thus be expressed as:

NPV =
$$\left[\frac{(1+i)^{n}-1}{i(1+i)^{n}}\right] - I$$
 (9-12)



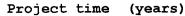


Figure 9-2. Net present value (NVP) of a project.

The internal rate of return (IRR) is that value of i that makes NPV equal to 0. Therefore, if NPV is set to 0, the IRR that makes the future cash flows equal to the investment (the "break-even" point) can be estimated. Equation 9-12 then becomes:

$$\frac{I}{C} = \frac{(1+i)^n - 1}{i(1+i)^n}$$
(9-13)

where I = investment

C = cash flow for each year

i = rate of return (IRR/100)

n = years of project life

Equation 9-13 is defined as the annuity with I, the present value, and C, the equal payments, over n, years at interest, i. The I/C term is referred to as the payback period. If I and C are evaluated and the life of the project is known, i can be computed by trial and error. The value of i can be determined from Figure 9-3.

Rate of	1%	2%	3%	4%	5%	6%	7%	88	98	10%
discount										
Year										
0	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000
1	0.9901	0.9804	0.9709	0.9615	0.9524	0.9434	0.9346	0.9259	0.9174	0.9091
2	0.9803	0.9612	0.9426	0.9246	0.9070	0.8900	0.8734	0.8573	0.8417	0.8264
3	0.9706	0.9423	0.9151	0.8890	0.8638	0.8396	0.8163	0.7938	0.7722	0.7513
4	0.9610	0.9238	0.8885	0.8548	0.8227	0.7921	0.7629	0.7350	0.7084	0.6830
5	0.9515	0.9057	0.8626	0.8219	0.7835	0.7473	0.7130	0.6806	0.6499	0.6209
6	0.9420	0.8880	0.8375	0.7903	0.7462	0.7050	0.6663	0.6302	0.5963	0.5645
7	0.9327	0.8706	0.8131	0.7599	0.7107	0.6651	0.6227	0.5835	0.5470	0.5132
8	0.9235	0.8535	0.7894	0.7307	0.6768	0.6274	0.5820	0.5403	0.5019	0.4665
9	0.9143	0.8368	0.7664	0.7026	0.6446	0.5919	0.5439	0.5002	0.4604	0.4241
10	0.9053	0.8203	0.7441	0.6756	0.6139	0.5584	0.5083	0.4632	0.4224	0.3855
11	0.8963	0.8043	0.7224	0.6496	0.5847	0.5268	0.4751	0.4289	0.3875	0.3505
12	0.8874	0.7885	0.7014	0.6246	0.5568	0.4970	0.4440	0.3971	0.3555	0.3186
13	0.8787	0.7730	0.6810	0.6006	0.5303	0.4688	0.4150	0.3677	0.3262	0.2897
14	0.8700	0.7579	0.6611	0.5775	0.5051	0.4423	0.3878	0.3405	0.2992	0.2633
15	0.8613	0.7430	0.6419	0.5553	0.4810	0.4173	0.3624	0.3152	0.2745	0.2394
16	0.8528	0.7284	0.6232	0.5339	0.4581	0.3936	0.3387	0.2919	0.2519	0.2176
17	0.8444	0.7142	0.6050	0.5134	0.4363	0.3714	0.3166	0.2703	0.2311	0.1978
18	0.8360	0.7002	0.5874	0.4936	0.4155	0.3503	0.2959	0.2502	0.2120	0.1799
19	0.8277	0.6864	0.5703	0.4746	0.3957	0.3305	0.2765	0.2317	0.1945	0.1635
20	0.8195	0.6730	0.5537	0.4564	0.3769	0.3118	0.2584	0.2145	0.1784	0.1486

 Table 9-2

 Discount Factors. This Table Shows the Present Value of Unity Discounted for Different Numbers of Years and at Different Rates of Discount

 Table 9-3

 Discount Factors. This Table Shows the Present Value of Unity Discounted for Different Numbers of Years and at Different Rates of Discount

Rate of	11%	12%	13%	14%	15%	16%	17%	18%	19%	20%
discount										
Year										
0	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000
1	0.9009	0.8929	0.8850	0.8772	0.8696	0.8621	0.8547	0.8475	0.8403	0.8333
2	0.8116	0.7972	0.7831	0.7695	0.7561	0.7432	0.7305	0.7182	0.7062	0.6944
3	0.7312	0.7118	0.6931	0.6750	0.6575	0.6407	0.6244	0.6086	0.5934	0.5787
4	0.6587	0.6355	0.6133	0.5921	0.5718	0.5523	0.5337	0.5158	0.4987	0.4823
5	0.5935	0.5674	0.5428	0.5194	0.4972	0.4761	0.4561	0.4371	0.4190	0.4019
6	0.5346	0.5066	0.4803	0.4556	0.4323	0.4104	0.3898	0.3704	0.3521	0.3349
7	0.4817	0.4523	0.4251	0.3996	0.3759	0.3538	0.3332	0.3139	0.2959	0.2791
8	0.4339	0.4039	0.3762	0.3506	0.3269	0.3050	0.2848	0.2660	0.2487	0.2326
9	0.3909	0.3606	0.3329	0.3075	0.2843	0.2630	0.2434	0.2255	0.2090	0.1938
10	0.3522	0.3220	0.2946	0.2697	0.2472	0.2267	0.2080	0.1911	0.1756	0.1615
11	0.3173	0.2875	0.2607	0.2366	0.2149	0.1954	0.1778	0.1619	0.1476	0.1346
12	0.2858	0.2567	0.2307	0.2076	0.1869	0.1685	0.1520	0.1372	0.1240	0.1122
13	0.2575	0.2292	0.2042	0.1821	0.1625	0.1452	0.1299	0.1163	0.1042	0.0935
14	0.2320	0.2046	0.1807	0.1597	0.1413	0.1252	0.1110	0.0985	0.0876	0.0779
15	0.2090	0.1827	0.1599	0.1401	0.1229	0.1079	0.0949	0.0835	0.0736	0.0649
16	0.1883	0.1631	0.1415	0.1229	0.1069	0.0930	0.0811	0.0708	0.0618	0.0541
17	0.1696	0.1456	0.1252	0.1078	0.0929	0.0802	0.0693	0.0600	0.0520	0.0451
18	0.1528	0.1300	0.1108	0.0946	0.0808	0.0691	0.0592	0.0508	0.0437	0.0376
19	0.1377	0.1161	0.0981	0.0829	0.0703	0.0596	0.0506	0.0431	0.0367	0.0313
20	0.1240	0.1037	0.0868	0.0728	0.0611	0.0514	0.0433	0.0365	0.0308	0.0261

Table 9-4
Discount Factors. This Table Shows the Present Value of Unity Discounted
for Different Numbers of Years and at Different Rates of Discount

Rate of discount	21%	22%	23%	24%	25%	26%	278	28%	29%	30%
Year										
0	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000
1	0.8264	0.8197	0.8130	0.8065	0.8000	0.7937	0.7874	0.7813	0.7752	0.7692
2	0.6830	0.6719	0.6610	0.6504	0.6400	0.6299	0.6200	0.6104	0.6009	0.5917
3	0.5645	0.5507	0.5374	0.5245	0.5120	0.4999	0.4882	0.4768	0.4658	0.4552
4	0.4665	0.4514	0.4369	0.4230	0.4096	0.3968	0.3844	0.3725	0.3611	0.3501
5	0.3855	0.3700	0.3552	0.3411	0.3277	0.3149	0.3027	0.2910	0.2799	0.2693
6	0.3186	0.3033	0.2888	0.2751	0.2621	0.2499	0.2383	0.2274	0.2170	0.2072
7	0.2633	0.2486	0.2348	0.2218	0.2097	0.1983	0.1877	0.1776	0.1682	0.1594
8	0.2176	0.2038	0.1909	0.1789	0.1678	0.1574	0.1478	0.1388	0.1304	0.1226
9	0.1799	0.1670	0.1552	0.1443	0.1342	0.1249	0.1164	0.1084	0.1011	0.0943
10	0.1486	0.1369	0.1262	0.1164	0.1074	0.0992	0.0916	0.0847	0.0784	0.0725
11	0.1228	0.1122	0.1026	0.0938	0.0859	0.0787	0.0721	0.0662	0.0607	0.0558
12	0.1015	0.0920	0.0834	0.0757	0.0687	0.0625	0.0568	0.0517	0.0471	0.0429
13	0.0839	0.0754	0.0678	0.0610	0.0550	0.0496	0.0447	0.0404	0.0365	0.0330
14	0.0693	0.0618	0.0551	0.0492	0.0440	0.0393	0.0352	0.0316	0.0283	0.0254
15	0.0573	0.0507	0.0448	0.0397	0.0352	0.0312	0.0277	0.0247	0.0219	0.0195
16	0.0474	0.0415	0.0364	0.0320	0.0281	0.0248	0.0218	0.0193	0.0170	0.0150
17	0.0391	0.0340	0.0296	0.0258	0.0225	0.0197	0.0172	0.0150	0.0132	0.0116
18	0.0323	0.0279	0.0241	0.0208	0.0180	0.0156	0.0135	0.0118	0.0102	0.0089
19	0.0267	0.0229	0.0196	0.0168	0.0144	0.0124	0.0107	0.0092	0.0079	0.0068
20	0.0221	0.0187	0.0159	0.0135	0.0115	0.0098	0.0084	0.0072	0.0061	0.0053

 Table 9-5

 Discount Factors. This Table Shows the Present Value of Unity Discounted for Different Numbers of Years and at Different Rates of Discount

Rate of	31%	32%	33%	34%	35%	36%	37%	38%	39%	40%
discount										
Year									<u> </u>	
0	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000
1	0.7634	0.7576	0.7519	0.7463	0.7407	0.7353	0.7299	0.7246	0.7194	0.7143
2	0.5827	0.5739	0.5653	0.5569	0.5487	0.5407	0.5328	0.5251	0.5176	0.5102
3	0.4448	0.4348	0.4251	0.4156	0.4064	0.3975	0.3889	0.3805	0.3724	0.3644
4	0.3396	0.3294	0.3196	0.3102	0.3011	0.2923	0.2839	0.2757	0.2679	0.2603
5	0.2592	0.2495	0.2403	0.2315	0.2230	0.2149	0.2072	0.1998	0.1927	0.1859
6	0.1979	0.1890	0.1807	0.1727	0.1652	0.1580	0.1512	0.1448	0.1386	0.1328
7	0.1510	0.1432	0.1358	0.1289	0.1224	0.1162	0.1104	0.1049	0.0997	0.0949
8	0.1153	0.1085	0.1021	0.0962	0.0906	0.0854	0.0806	0.0760	0.0718	0.0678
9	0.0880	0.0822	0.0768	0.0718	0.0671	0.0628	0.0588	0.0551	0.0516	0.0484
10	0.0672	0.0623	0.0577	0.0536	0.0497	0.0462	0.0429	0.0399	0.0371	0.0346
11	0.0513	0.0472	0.0434	0.0400	0.0368	0.0340	0.0313	0.0289	0.0267	0.0247
12	0.0392	0.0357	0.0326	0.0298	0.0273	0.0250	0.0229	0.0210	0.0192	0.0176
13	0.0299	0.0271	0.0245	0.0223	0.0202	0.0184	0.0167	0.0152	0.0138	0.0126
14	0.0228	0.0205	0.0185	0.0166	0.0150	0.0135	0.0122	0.0110	0.0099	0.0090
15	0.0174	0.0155	0.0139	0.0124	0.0111	0.0099	0.0089	0.0080	0.0072	0.0064
16	0.0133	0.0118	0.0104	0.0093	0.0082	0.0073	0.0065	0.0058	0.0051	0.0046
17	0.0101	0.0089	0.0078	0.0069	0.0061	0.0054	0.0047	0.0042	0.0037	0.0033
18	0.0077	0.0068	0.0059	0.0052	0.0045	0.0039	0.0035	0.0030	0.0027	0.0023
19	0.0059	0.0051	0.0044	0.0038	0.0033	0.0029	0.0025	0.0022	0.0019	0.0017
20	0.0045	0.0039	0.0033	0.0029	0.0025	0.0021	0.0018	0.0016	0.0014	0.0012

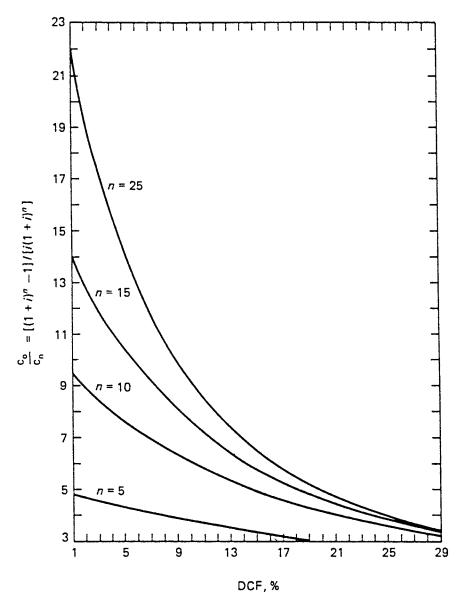


Figure 9-3. Graphical solution for simplified DCF calculation. (Constant cash flow, compounded annually, no salvage or working capital return.) Source: Horwitz [1].

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(text continued from page 729)

The NPV measures the direct incentive to invest in a given proposal as a bonus or premium over the amount an investor could otherwise earn by investing the same money in a safe alternative, which would yield a return calculated at the rate, i. The resulting NPV from a project's cash flow is a measure of the cash profit that the project will produce after recovering the initial investment and meeting all costs, including the cost of capital. The more positive the NPV is, the more attractive the proposition. If NPV is 0, the viability of the project is marginal; if it is negative, the proposal is unattractive.

Discounted Cash Flow Rate of Return (DCFRR)

The discounted cash flow rate of return is known by other terms, for example, the profitability index, the true rate of return, the investor's rate of return, and the internal rate of return. It is defined as the discount rate, i, which makes the NPV of a project equal to zero. This can be expressed mathematically by

NVP =
$$\sum_{t=0}^{t=n} \frac{C_t}{(1+t)^t} = 0$$
 (9-14)

The main advantage of DCFRR over NPV is that it is independent of the zero or base year that is chosen. In contrast, the value of NPV varies acccording to the zero year chosen. In calculating the NPV, the cost of capital has to be explicitly included as i in the discounting calculations. In computing the DCFRR, the cost of capital is not included. Instead the value calculated for i is compared with the cost of capital to see whether the project is profitable. The DCFRR for a project is the rate of return on investment. It measures the efficiency of the capital and determines the earning power of the project investment. Therefore, a DCFRR of say 20 percent implies that 20 percent per year will be earned on the investment, in addition to which the project generates sufficient money to repay the original investment plus any interest payable on borrowed capital plus all taxes and expenses.

Net Return Rate (NRR)

The net return rate is analogous to the rate of return and is the net average discounted "return" on the investment over and above the cost of capital. This is defined by: 736 Fortran Programs for Chemical Process Design

$$NRR = \frac{NPV}{\left(\frac{Discounted}{investment}\right)\left(\frac{Project}{life}\right)} \times 100$$
(9-15)

where the investment is discounted to the same point as the NPV. Holland *et al.* [7] introduced the NPV/investment ratio as a normalized measure of the total discounted return over the life of the investment. Ward [4] showed that the NPV can be divided by the number of cash flow increments (venture lifetime) so that the NRR corresponds to the average discounted net return on investment. The cost of capital is already accounted for by the discount rate in the NPV computation, and therefore the NRR is the true return rate.

Depreciation

Estimation of depreciation charges may be based on:

- 1. cost of operation
- 2. tax allowance
- 3. means of building up a fund to finance plant replacement
- 4. measure of falling value

The annual depreciation charge can be calculated using "straight line" depreciation, and is expressed as:

$$D = \frac{C_{FC} - S}{n}$$
(9-16)

where D = annual depreciation

 C_{FC} = initial fixed capital cost

n = n years of projected life

S = salvage value

Assuming that C_{FC} is the initial fixed capital investment, and S is the projected salvage value at the end of n years of projected life, then the depreciated rate, d_i , for any particular year, j, is:

$$D_{j} = (C_{FC} - S)d_{j}$$
 (9-17)

where D_i = annual depreciation charge

With the straight line calculation procedure, where d_j is constant, combining Equation 9-16 and Equation 9-17 gives:

$$d = \frac{1}{n} \tag{9-18}$$

The various components of a plant such as equipment, buildings, and improvements are characterized by projected lifetimes. During this period, each item depreciates from its initial investment cost, C_{FC} , to a salvage value, S, over the period of n years of its projected lifetime. At the end of any particular year, k, the depreciated value or book value, V_k , is:

$$\mathbf{V}_{k} = \mathbf{C}_{FC} - \sum_{i}^{k} \mathbf{D}_{j}$$
(9-19)

where D_j is the annual depreciation charge for year j. Substituting Equation 9-17 into Equation 9-19 gives

$$V_{k} = C_{FC} - \sum_{i}^{k} (C_{FC} - S)d_{j}$$

= $C_{FC} - (C_{FC} - S)\sum_{i}^{k} d_{j}$ (9-20)

For the straight line depreciation procedure,

$$\sum_{i}^{k} d_{j} = \sum_{i}^{k} d = kd = \frac{1}{n}$$
(9-21)

and

$$V_{k} = C_{FC} - \frac{k}{n}(C_{FC} - S)$$
 (9-22)

Double Declining Balance (DDB) Depreciation

Equipment and complete plants depreciate and lose value more rapidly in the early stages of life. The depreciation based upon the declining book value balance can be expressed as:

$$D_j = d_j \cdot V_{j-1}$$
 (9-23)

The rate of depreciation, d_j , is the same for each year, j; however, the depreciation charges decrease each year because the book value decreases each year. For a declining balance method, the depreciation

rate of decline is up to twice, but no more than twice the straight line rate. This is given by:

$$d_j = d = \frac{2}{n} \tag{9-24}$$

Capitalized Cost

The capitalized cost, C_K , of a piece of equipment of a fixed capital cost, C_{FC} , having a finite life of n years and an annual interest rate, i, is defined by

$$(C_{K} - C_{FC})(1+i)^{n} = C_{K} - S$$
 (9-25)

where S = salvage or scrap value

 C_{K} is in excess of C_{FC} by an amount, which, when compounded at an annual interest rate, i, for n years, will have a future worth of C_{K} less the salvage or scrap value, S. If the renewal cost of the equipment and the interest rate are constants at $(C_{FC} - S)$ and i, then C_{K} is the amount of the capital required to replace the equipment in perpetuity.

Rearranging Equation 9-25 gives

$$C_{K} = \left[C_{FC} - \frac{S}{(1+i)^{n}}\right] \left[\frac{(1+i)^{n}}{(1+i)^{n} - 1}\right]$$
(9-26)

or

$$C_{K} = (C_{FC} - S.f_{d})f_{K}$$
 (9-27)

where $f_d = discount factor$

 f_{K} = the capitalized cost factor = $(1 + i)^{n}/[(1 + i)^{n} - 1]$

The Average Rate Of Return (ARR)

The average rate of return (ARR) method averages out the cash flow over the life of the project. This is defined by

$$ARR = \frac{\text{average cash flow}}{\text{original investment}} \times 100$$
(9-28)

The higher the percentage value of the ARR, the better the profitability of the project.

Present Value Ratio (Present Worth Ratio)

A commonly used profitability index in conjunction with the NPV method shows how closely a project has met the criterion of economic performance. This index is known as the present value ratio (PVR) or present worth ratio (PWR), and is defined as

$$\frac{\text{Present}}{\text{value}} = \frac{\text{present value of all positive cash flows}}{\text{present value of all negative cash flows}}$$
(9-29)

The present value ratio (PVR) gives an indication of how much the project makes relative to the investment. A ratio of 1.0 shows that the income just matches the expected income from capital invested for a given interest rate. A ratio of less than 1.0 indicates that the income does not come up to the minimum expectations. A ratio of more than 1.0 means that the project exceeds the minimum expectations.

Profitability

A project is profitable if its earnings are greater than the cost of capital. In addition, the larger the additional earnings, the more profitable the venture, and the greater the justification for putting the capital at risk. Therefore, a profitability estimate attempts to quantify the risk taken. The methods used to assess profitability are:

- 1. return on investment (ROI)
- 2. payback period (PB)
- 3. net present value (NPV)
- 4. discounted cash flow rate of return (DCFRR)
- 5. net return rate (NRR)
- 6. equivalent maximum investment period (EMIP)
- 7. interest recovery period (IRP)
- 8. rate of return on depreciated investment
- 9. rate of return on average investment
- 10. capitalized cost
- 11. average rate of return (ARR)
- 12. present value ratio (PVR)

Abrams [11] has listed other methods for assessing a project's profitability.

Economic Analysis

Computer programs have been developed to estimate the net present value (NPV), present value ratio (PVR), net return rate (NRR), average rate of return (ARR), payback period (PBP) and discounted cash flow rate of return (DCFRR). These analyses are done for a given cash flow over the operating life of a project. These programs can be incorporated as subroutines into larger programs, if required. In addition, a detailed computer program has been developed to review an economic project using Kirkpatrick's [12] input data. These data are defined as follows:

Annual Revenue, \$. The money received (sales minus the cost of sales) for one-year production from the plant. This is assumed as being constant for life of the project.

Annual Operating Cost, **\$.** The cost of raw materials, labor, utilities, administration, insurance, and royalties, etc., but does not include debt service payments.

Depreciating Base, \$. The capitalized cost of the facility and less non-depreciable items, such as land and inventory. No salvage is sub-tracted because double declining balance depreciation is used.

Project Life, Years. The length of time for which the facility is to be operated. It is also the term of the loan and the depreciation time.

Initial Loan, \$. The capitalized cost minus owner equity.

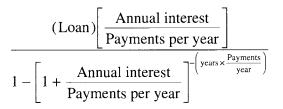
Payments/Year. The number of payments on the loan per year: 1-annually, 2-biannually, 4-quarterly, and 12-monthly.

Periodic Interest Rate. The annual interest rate divided by the payments per year. For a 10 percent annual interest rate and monthly repayments, this is 0.10/12.

Investment Tax Credit, \$. The percentage of the initial investment allowed as a tax credit in the year the investment is made.

Tax Rate. The percentage tax that must be paid on the project's pretax income. This rate is assumed to remain constant during the life of the project.

Debt Service/Period, \$. The amount of each loan payment, that is:



The Salvage Value (or Scrap Value). The projected salvage value of equipment at the end of the project lifetime is that portion of the fixed capital that cannot be depreciated.

Land Value. Land is not considered depreciable. It can be used indefinitely for succeeding projects on a specific site, or it can be sold.

Working capital. The capital invested in various necessary inventoried items, which are completely recoverable at any time.

The calculations for every year of the project life are:

- 1. Calculate the depreciation as double declining (that is, twice the depreciation based divided by project life), and subtract it from the depreciation base.
- 2. Calculate the yearly interest and principal of the debt service payments.
- 3. Subtract from the revenue, the operating cost, annual depreciation, and interest.
- 4. Subtract from positive pre-tax income, until no negative pre-tax income remains, if the pre-tax income is negative or has been negative in a prior year.
- 5. Calculate the income tax due at the given rate, and apply the investment tax credit against the income tax due until the credit is exhausted, if after Step 4, the pre-tax income is still positive.
- 6. Deduct the income tax remaining after Step 5 from the pre-tax income, leaving the after-tax income.
- 7. Add the after-tax income to the depreciation for the year, yielding the cash flow.
- 8. Determine the loan balance at the end of the year.
- 9. Determine the amount of unused depreciation after the last year of the project's life. Because double declining depreciation does not totally exhaust the depreciation account, the unused depreciaton should be added to the cash flow of the last year as salvage value recovered at the termination of the project under consideration.

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- 10. Calculate the net present value, NPV, for a known discount rate.
- 11. Determine the present value ratio, PVR.
- 12. Calculate the net return rate, NRR.
- 13. Calculate the average rate of return, ARR.
- 14. Estimate the payback period, PBP.

For a given discount rate, calculate the discount factor during the operating life of the project. Multiply the discount factor by the cash flow and obtain the net present value and the net return rate. Valle-Riesta [8] has listed alternative methods of calculating cash flow, as shown in Table 9-6.

Methods 1 and 2 in Table 9-6 give identical results.

Method 1	Method 2
1. Cost of Manufacture (COM) includes depreciation	1. Cost of Manufacture (COM) includes depreciation
2. General expenses	2. Depreciation
3. Total operating expense (3) = $(1) + (2)$	3. COM less depreciation (3) + (1) - (2)
4. Total sales	4. General expenses
5. Profit before taxes	5. Out of pocket expenditure
6. Income taxes (6) $\approx 1/2$ (5)	6. Total sales
7. Profit after tax (7) = $(5) - (6) \approx 1/2$ (5)	7. Cash flow before taxes (7) = (6) - (5)
8. Depreciation	8. Profit before taxes (8) = $(7) - (2)$
9. Continuous cash flow from project (9) = (7) + (8)	9. Income tax (9) $\approx 1/2$ (8)
	10. Profit after tax (10) = (8) - (9) $\approx 1/2$ (8)
	11. Continuous cash flow from project (11) = $(10) + (2)$

Table 9-6 Method of Computing Cash Flow from Project

Source: Valle-Riesta [8]

Cash flows into a project can be time-dependent, in which the cash flows occur in a continuous process rather than on a one-time basis. Figure 9-4 illustrates a continuous cash flow diagram. Continuous cash flows into the project are from sales revenues, and cash flows out are for out-of-pocket expenses. The difference between the incoming and outgoing cash flow is the net cash flow generated by the project. The combined net cash flow generated by all the company's projects is reduced by the income tax payment to yield the net continuous cash flow. Expressing the definition of profit with the continuous cash flow results in the following:

Profit before taxes = continuous cash flow – depreciation before taxes

Profit after taxes = continuous cash flow – depreciation after taxes

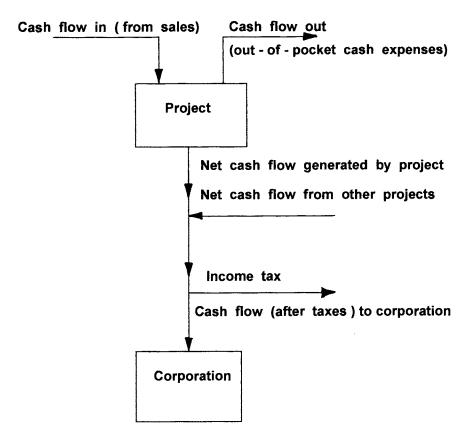


Figure 9-4. A continuous cash flow diagram. Source: Valle-Riestra [8].

It is useful to keep in mind that:

Cash flow \neq profit

Allen [9] has provided a systematic procedure for assessing investment proposals for new plant and equipment, exploiting new technology, and replacing uneconomic, inefficient, and obsolete plants, process, and equipment.

Inflation

A decrease in the average purchasing value of currency is referred to as inflation. An inflation rate of 15 percent, for example, means that the average cost of goods and services will increase 15 percent in one year. The result is that commencing construction one year early will reduce the amount of money expended by 15 percent. Since the 1980s, inflation has been considered in most economic project evaluations.

When inflation is used in economic evaluations, all items except interest on a loan and depreciation are considered to increase in value at the same rate as inflation. Generally, interest is set at the time a loan is negotiated and does not change with inflation. In addition, depreciation depends on the method (for example, straight line or double declining balance) used and the capital charges incurred before start-up are not affected by the inflation rate after start-up. Determining the profitability of a project (for example, NPV), the interest rate is assumed to be greater than the inflation rate. Money may be lost on the project while the net present value indicates the opposite, if the inflation rate is greater than the interest rate. There are cases in which the interest rate is set at the expected inflation rate plus a real expected interest rate. The real expected interest rate is the interest rate that is used to calculate the net present value when there is no inflation. Alternatively, the present value is calculated using the inflation rate as the interest rate, the net present value is then determined using the real expected interest rate.

CAPITAL COST ESTIMATION

Capital cost estimation is an essential part of investment appraisal. Many types of capital cost estimates are made, ranging from the order of magnitude to detailed estimates requiring the collection of accurate technical data. The American Association of Cost Engineers defined five types of cost estimates as follows: **Order-of-magnitude estimate (ratio estimate).** Approximate method based on cost data for previous similar types of plant. Probable error within 10 to 50 percent.

Study estimate (factored estimate). Better than order-of-magnitude but requires knowledge of major items of equipment. Used for feasibility surveys. Probable error up to 30 percent.

Preliminary estimate (budget authorization estimate). More detailed information required than for study estimate. Probable error up to 20 percent.

Definitive estimate (project control estimate). Based on considerable data obtained before preparing completed drawings and specifications. Probable error within 10 percent.

Detailed estimate (firm or contractor's estimate). Requires completed drawings and specifications. Probable error within 10 percent.

Process plant designs start from preliminary designs based on approximate technical data, calculations, and cost data to final designs that require detailed and accurate data, calculations, and quotations. Cost estimates of a proposed plant are continuously carried out during the development of a process from the laboratory to construction.

The total capital cost, C_{TC} , of a project consists of the fixed capital cost, C_{FC} , plus the working capital, C_{WC} , plus the cost of land and any other non-depreciable assets, C_L . This is given by

$$C_{TC} = C_{FC} + C_{WC} + C_L$$
 (9-30)

The fixed capital cost, C_{FC} , is the capital required to provide all the depreciable facilities. C_{FC} may be divided into two classes known as the battery limits and auxiliary facilities. The boundary of battery limits includes all manufacturing and processing equipment. The auxiliary facilities are the storage areas, administration offices, utilities and other essential and non-essential supporting facilities.

Equipment Cost Estimations By Capacity Ratio Exponents

It is often necessary to calculate the cost of a piece of equipment when there are no available cost data for the particular size of capacity. If the cost of a piece of equipment or plant size or capacity, Q_1 , is C_1 , the cost C_2 of a similar piece of equipment or plant size or capacity, Q_2 , can be calculated from the equation

$$\mathbf{C}_2 = \mathbf{C}_1 \left(\frac{\mathbf{Q}_2}{\mathbf{Q}_1}\right)^{\mathrm{m}} \tag{9-31}$$

where

the value, m, depends on the type of equipment or plant. The value of the index, m, is generally taken as 0.6 [13], the well known six-tenths rule. This value can be used to get a rough estimate of the capital cost, if there are insufficient data to calculate the index for the particular size of equipment required. Table 9-7 lists values of m for various types of equipment. Cost indexes should be used to bring the cost data to a desired year. A cost index is a value for a given point in time showing the cost at that time relative to a certain base time. If the cost at some time in the past is known, the equipment cost at the present time can be determined from the equation

Present cost = previous cost
$$\left(\frac{\text{present index}}{\text{previous index}}\right)$$
 (9-32)

Cost indexes are used to give a general estimate, but no index can account for all the factors. Many different types of cost indexes are

Equipment	Exponent (m)
Reciprocating compressor	0.75
Turboblowers compressor	0.5
Electric motors	0.8
Evaporators	0.5
Heat exchangers	0.65 to 0.95
Piping	0.7 to 0.9
Pumps	0.7 to 0.9
Rectangular tanks	0.5
Spherical tanks	0.7
Towers, constant diameter	0.7
Towers, constant height	1.0

 Table 9-7

 Typical Exponents for Equipment Cost vs Capacity

Source: Institution of Chemical Engineers [15].

published in magazines such as the *Chemical Engineering Plant* cost index, *Engineering News-Records* construction index, the *Oil & Gas Journal*, and *Process Engineering*. Table 9-8 shows the international plant cost indexes from March 1992 to February 1993.

Factored Cost Estimate

The purchased cost of an item of equipment, free on board (FOB) is quoted by a supplier, and may be multiplied by a factor of 1.1 to give the approximate delivered cost. The factorial methods for estimating the total installed cost of a process plant are based on a combination of materials, labor, and overhead cost components. The fixed capital cost, C_{CF} , of a plant based on design can be estimated using the Lang Factor method [14] given by the equation

$$C_{FC} = f_L \sum C_{EQ}$$
(9-33)

where $f_L = 3.10$ for solids processing

 $f_L = 3.63$ for mixed solids-fluid processing

 $f_L = 4.74$ for fluid processing

 ΣC_{EQ} is the sum of the delivered costs of all the major items of process equipment. The major advantage of the Lang method is that the cost of equipment is available.

Other methods for estimating capital investment consider the fixedcapital investment required as a separate unit. These are known as the functional-unit estimates, the process step scoring method, and the modular estimate.

The functional unit may be characterized as a unit operation, unit process, or separation method that involves energy transfer, moving parts, or a high level of internals. The unit includes all process streams together with side or recycle streams. Bridgwater [16] proposed seven functional units, namely, compressor, reactor, absorber, solvent extractor, solvent recovery column, main distillation column, and furnace and waste heat boiler. Taylor [17] developed the step counting method, based on a system in which a complexity score accounting for factors such as throughput, corrosion problems, and reaction time is estimated for each process step. The modular estimate considers individual modules in the total system with each module consisting of a group of similar items. For the modular estimate, all heat exchangers are classified in one module, all furnaces in another, all vertical process vessels in another,

Table 9-8
International Plant Cost Indexes
Mar 1992-Feb 1993 Base Date Jan 1990 = 100

COUNTRY	Mar	Apr	May	Jun	Jul	Aug	Sep	Oct	Nov	Dec	Jan	Feb
Australia	107.7	107.9	107.9	108.2	108.4	108.6	108.7	108.8	108.8	109.0	109.3	109.4
Belgium	108.6	109.2	109.3	109.2	110.4	110.8 ^R	110.8	111.4	111.2	111.2	111.2	111.2
Canada	107.0	107.0	106.9	106.3	105.7		106.9	107.5	107.5	108.7	108.8	108.8
Denmark	107.0	108.4	108.5	109.3	110.4	108.0	108.5	108.9	108.5	110.4	110.7	-
France	106.5	106.9	107.0	107.1	107.5	107.5	108.1	108.3	108.5	108.6	109.2	109.5
Germany	113.8	114.0	114.0	114.9	115.0	115.0	115.4	115.4	115.4 ^F	116.2	116.3	116.4
Italy	109.5	109.6	109.6	109.9	110.1	110.0	110.2	110.5	110.5	110.7	-	-
Japan	103.0	102.8	102.8	110.4	102.7	101.3	103.1	103.3	103.4	102.5	100.3	103.0
Netherlands	106.8	108.0	108.1	108.1	108.8	108.8	108.8	108.7	108.7	108.7	109.5	109.5
New Zealand	103.9	104.4	104.4	104.4	105.0	105.0	105.0	105.3	105.4	105.4	-	_
Spain	111.1	113.2	113.2	113.3	114.4	114.4	114.4	114.7	114.8	114.8	-	-
Sweden	106.2	107.9	108.8	109.2	109.6	107.9	108.4	109.0	109.5	111.3	-	-
USA	101.7	101.1	101.6	101.4	100.4	100.8	100.7	100.9	100.5	100.7	100.7	100.7
UK	117.2	115.0	115.1	117.0	116.8	117.1	118.2	116.2	114.4	114.6	114.9	115.8

R = revised value, P = provisional value, F = forecast value

SOURCE: PROCESS ENGINEERING JUNE 1993

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etc. The total cost estimate is considered under six general groupings. These are chemical processing, solids handling, site development, industrial buildings, offsite facilities, and project indirects [18].

Nomenclature

```
ARR = average rate of return
```

C = cash flow for each year

 C_{CF} = net annual cash flow

- C_{FC} = fixed capital cost
- C_{K} = capitalized cost
- $C_L = land cost$

 C_0 = initial investment cost

 $C_n = \text{cash flow}$

 C_{TC} = total capital cost

 C_{wc} = working capital cost

 C_1 = capital cost of the designed plant

 C_2 = capital cost of the existing plant

D = annual depreciation

DCFRR = discounted cash flow rate of return

```
D_i = annual depreciation charge
```

 $d_i =$ depreciation rate

EMIP = equivalent maximum investment period

```
f_d = discount factor
```

 f_{κ} = capitalized cost factor

I = investment cost

IRP = interest recovery period

- IRR = internal rate of return
 - i = interest rate of return (ROI/100)
 - m = exponential power for cost capacity relationships

NPV = net present value

NRR = net return rate

n = years of project life

- PBP = payback period
- PV = present value
- PVR = present value ratio

PWR = present worth ratio

- Q_1 = capacity of the designed plant
- Q_2 = capacity of existing plant
- ROI = return of investment

S = salvage value

PROBLEMS AND SOLUTIONS

Problem 9-1

Consider a plant costing \$1,000,000 to build that produces a product A. For the same capital outlay a different plant can be erected to produce an alternative product B. Conditions are such that each plant will only be in operation for eight years and then both will be scrapped. The cash flows in each of the eight years obtained by selling A and B are shown in Table 9-9. A choice is being made about which plant is more profitable.

Solution

The computer program PROG91 calculates the net present value and the net return rate at discount rates of 5, 10, 15, 20, 25, 30, 35, and 40%. Tables 9-10 and 9-11 give both the input data and computer outputs for products A and B. Figure 9-5 shows a plot of the cumulative cash flows against the project life, and Figure 9-6 gives a plot of the NPV against the discount rate. Figure 9-5 gives the payback period of product A between 3 and 4 years and product B is about 2 years. The results show

Year	Product A Cash Flows \$	Product A Cumulative Cash Flows \$	Product B Cash Flows \$	Product B Cumulative Cash Flows \$
	-1,000,000	-1,000,000	-1,000,000	-1,000,000
1	100,000	-900,000	800,000	-200,000
2	200,000	-700,000	700,000	500,000
3	300,000	-400,000	600,000	1,100,000
4	400,000	0	500,000	1,600,000
5	500,000	500,000	400,000	2,000,000
6	600,000	1,100,000	300,000	2,300,000
7	700,000	1,800,000	200,000	2,500,000
8	800,000	2,600,000	100,000	2,600,000

Table 9-9Annual Cash Flows and Cumulative Cash Flows

Table 9-10 Input Data and Computer Output for the Net Present Value of Product A at 10 Percent Discount Rate

DATA91.DAT				<u> </u>
8 10 -1000000 200000 300000 400000 500000 600000 700000 800000				
*****		ESENT VALUE CALCULA		*****
	9 YEARI	LY CASH FLOWS INCLU	DING YEAR O	
	10.00	PERCENTAGE ANNUAL D	ISCOUNT RATE	
YEAR	CASH FLOWS (\$)	CUMULATIVE CASH FLOWS (\$)	DISCOUNT FACTOR	PRESENT VALUE (\$)
0	-1000000.00	-1000000.00	1.0000	-1000000.00
1	100000.00	-900000.00	.9091	90909.09
2 3	200000.00	-700000.00 -400000.00	.8264 .7513	165289.25
3 4	300000.00 400000.00	-400000.00	.6830	225394.43 273205.36
5	500000.00	500000.00	.6209	310460.63
6	600000.00	1100000.00	.5645	338684.31
7	700000.00	1800000.00	.5132	359210.63
8	800000.00	260000.00	.4665	373205.84
THE NET	PRESENT VALUE (\$)): 1136359	.53	
PRESENT	VALUE RATIO:	2.136		
THE NET	RETURN RATE:	14.20	\$	
THE AVE	RAGE RATE OF RETUR	RN: 45.00	*	
THE PAY	BACK PERIOD IS BE	TWEEN: 3 AN	ID 4 YEARS	

that product B gives greater present value ratios than product A. In addition, product B gives greater net present values at different discount rates than product A. These show that product B is more promising than product A. Further profitability indexes (ARR, NRR, PBP) suggest that product B is a better choice than product A.

Problem 9-2

Using the cash flows of products A and B in Problem 9-1, determine the discount cash flow rate of return (DCFRR).

Table 9-11 Input Data and Computer Output for the Net Present Value of Product B at 10 Percent Discount Rate

DATA91.DAT				
8 10 -1000000 800000 700000 600000 500000 400000 300000 200000 100000				
*****		ESENT VALUE CALCULAT		*****
	9 YEARI	LY CASH FLOWS INCLU	DING YEAR 0	
	10.00	PERCENTAGE ANNUAL D	ISCOUNT RATE	
YEAR	CASH FLOWS (\$)	CUMULATIVE CASH FLOWS (\$)	DISCOUNT FACTOR	PRESENT VALUE (\$)
0 1 2 3 4 5 6 7 8	$\begin{array}{c} -1000000.00\\ 80000.00\\ 700000.00\\ 50000.00\\ 500000.00\\ 400000.00\\ 30000.00\\ 20000.00\\ 100000.00\\ \end{array}$	-100000.00 -20000.00 50000.00 10000.00 200000.00 230000.00 250000.00 260000.00	1.0000 .9091 .8264 .7513 .6830 .6209 .5645 .5132 .4665	-1000000.00 727272.71 578512.37 450788.85 341506.70 248368.50 169342.16 102631.61 46650.73
PRESEN	F PRESENT VALUE (\$) F VALUE RATIO: F RETURN RATE:): 1665073. 2.665 20.81		
THE AVI	ERAGE RATE OF RETUR	RN: 45.00		

Solution

The computer program PROG92 determines the discounted cash flow rate of return (DCFRR). Tables 9-12 and 9-13 illustrate both the input data and computer outputs for products A and B. The DCFRR of product A is 28.264%, and that of product B is 64.863%. The results further confirm that product B is more promising than product A.

Problem 9-3

Calculate the yearly return of investment on the following financing data as shown in Table 9-14.

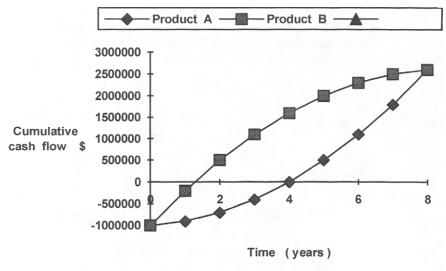


Figure 9-5. Cumulative cash flow diagram.

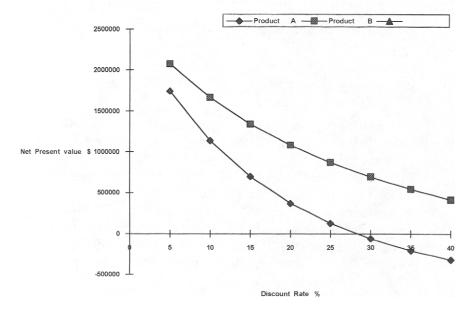


Figure 9-6. Net present value vs. discount rate.

Table 9-12 Input Data and Computer Output for the Discount Cash Flow Rate of Return (DCFRR) of Product A

9 -1000000 100000 200000 300000 400000 500000 600000 700000 800000			
*****		COUNT CASH FLOW RATE OF RETURN CALCULATION	****
	9 YEAR	YEARLY CASH FLOWS INCLUDING YEAR 0 CASH FLOW (\$)	

THE DISCOUNT CASH FLOW RATE OF RETURN (%): 28.264

Solution

The computer program PROG93 computes the following: the yearly cash flows, the cumulative cash flows, the present value, the net present value, present value ratio, average return rate, net return rate, and the payback period. Table 9-15 gives the input data and computer output for the cash flows during 10 years of the project life, and at a discount rate of 5 percent. The computed salvage value of the investment is \$4,294,967. The cash flows generated by the computer program PROG93 is then used to determine the DCFRR from program PROG92. Table 9-16 lists the input data and computer output for DCFRR of the investment as 6.347 percent.

Table 9-13 Input Data and Computer Output for the Discount Cash Flow Rate of Return (DCFRR) of Product B

DATA92.DAT 9 -1000000 800000 700000 600000 500000 400000 300000 200000 100000		
****	*******	SCOUNT CASH FLOW RATE OF RETURN CALCULATION
	YEAR	CASH FLOW (\$)
**********	******** 0 1 2 3 4 5 6 7 8	-100000.00 80000.00 70000.00 50000.00 50000.00 40000.00 30000.00 20000.00 100000.00

THE DISCOUNT CASH FLOW RATE OF RETURN (%): 64.863

Table 9-14 Financing Data

1.	Annual revenue, \$	9,000,000
2.	Annual operating cost \$	1,200,000
3.	Depreciating base, \$	40,000,000
4.	Project life, years	10
5.	Initial loan, \$	30,000,000
6.	Number of payment per year	12
7.	Annual interest, %	10
8.	Investment credit, %	10
9.	Tax rate, %	50
10.	Rate of return, %	5

Table 9-15 Input Data and Computer Output for the Economic Project Evaluation at 5 Percent Discount Rate

DATA93.DAT		
900000 1200000 40000000 10 30000000 12 10 50 50 5		
*****	ECONOMIC EVALUATION	OF A PROJECT **********
1	ANNUAL REVENUE \$:	9000000.00
2	ANNUAL OPERATING COST \$:	1200000.00
3	DEPRECIATION BASE \$:	4000000.00
4	PROJECT LIFE, YEARS:	10.
5	INITIAL LOAN \$:	3000000.00
6	NUMBER OF PAYMENTS PER YEAR	: 12.
7	ANNUAL INTEREST, %:	10.
8	INVESTMENT CREDIT, %:	10.
9	TAX RATE, %:	50.
10	DISCOUNT RATE %:	5.00

*

******	***************************************
REVENUE \$:	900000.00
OPERATING COST \$:	1200000.00
DEPRECIATION \$:	8000000.00
INTEREST \$:	2917171.30
PRE-TAX INCOME \$:	-3117171.30
TAX AT 50.%, \$:	.00
AFTER TAX INCOME \$:	-3117171.30
CASH FLOW \$:	4882828.70

YEAR 2	*****
REVENUE \$:	9000000.00
OPERATING COST \$:	1200000.00
DEPRECIATION \$:	6400000.00
INTEREST \$:	2724472.52
PRE-TAX INCOME \$:	-1324472.52
TAX AT 50.%, \$:	.00
AFTER TAX INCOME \$:	-1324472.52
CASH FLOW \$: ************************************	5075527 .48 *******
YEAR 3	*****
REVENUE \$:	9000000.00
OPERATING COST \$:	1200000.00
DEPRECIATION \$:	5120000.00
INTEREST \$:	2511595.66
PRE-TAX INCOME \$:	168404.34
TAX AT 50.%, \$:	.00
AFTER TAX INCOME \$:	168404.34
CASH FLOW \$: ************************************	5288404.3 4
YEAR 4	******
REVENUE \$:	9000000.00
OPERATING COST \$:	1200000.00
DEPRECIATION \$:	4096000.00
INTEREST \$:	2276427.81
PRE-TAX INCOME \$:	1427572.19
TAX AT 50.%, \$:	.00
AFTER TAX INCOME \$:	1427572.19
CASH FLOW \$:	5523572.19
	(table continued on next page)

Table 9-15 (continued)

	YEAR 5	****
REVENUE S:		900000.00
OPERATING COST \$:		1200000.00
DEPRECIATION \$:		3276800.00
INTEREST \$:		2016634.81
PRE-TAX INCOME \$:		2506565.19
TAX AT 50.%, \$:		.00
AFTER TAX INCOME \$:		2506565.19
CASH FLOW \$:	*****	5783365.19 ******

Table 9-15 (continued)

YEAR 6

A	9000000.00
REVENUE \$:	900000.00
OPERATING COST \$:	1200000.00
DEPRECIATION \$:	2621440.00
INTEREST \$:	1729638.09
PRE-TAX INCOME \$:	3109819.75
TAX AT 50.%, \$:	.00
AFTER TAX INCOME \$:	3109819.75
CASH FLOW \$:	5731259.75

REVENUE \$:	9000000.00
OPERATING COST \$:	1200000.00
DEPRECIATION \$:	2097152.00
INTEREST \$:	1412589.07
PRE-TAX INCOME \$:	4290259.00
TAX AT 50.%, \$:	.00
AFTER TAX INCOME \$:	4290259.00
CASH FLOW \$:	6387411.00

YEAR 8	****
REVENUE \$:	900000.00
OPERATING COST \$:	1200000.00
DEPRECIATION \$:	1677722.00
INTEREST \$:	1062340.87
PRE-TAX INCOME \$:	5059937.50
TAX AT 50.%, \$:	2230008.00
AFTER TAX INCOME \$:	2829929.50
CASH FLOW \$: ************************************	4507651.13 ******

Table 9-15 (continued)

.

YEAR	9 ********
REVENUE \$:	900000.00
OPERATING COST \$:	1200000.00
DEPRECIATION \$:	1342177.00
INTEREST \$:	675417.10
PRE-TAX INCOME \$:	5782405.50
TAX AT 50.%, \$:	2891202.75
AFTER TAX INCOME \$:	2891202.75
CASH FLOW \$:	4233380.00

YEAR 10 *****

REVENUE \$:	900000.00
OPERATING COST \$:	1200000.00
DEPRECIATION \$:	1073742.00
INTEREST \$:	247977.37
PRE-TAX INCOME \$:	6478281.00
TAX AT 50.%, \$:	3239140.50
AFTER TAX INCOME \$:	3239140.50
CASH FLOW \$:	4312882.38
SALVAGE VALUE \$:	4294967.30

(table continued on next page)

Table 9-15 (continued)

ECONOMIC PROJECT EVALUATION NET PRESENT VALUE AT A GIVEN DISCOUNT RATE				
*******		YEARLY CASH FLOWS	INCLUDING YEAR 0	*****
	5.00	PERCENTAGE ANNUA	AL DISCOUNT RATE	
	YEARLY CASH F	LOWS (\$)		
YEAR	CASH FLOWS (\$) CUMULATIVE CASH FLOWS (\$)	DISCOUNT) FACTOR	PRESENT VALUE (\$)
0	-40000000.00	-40000000.00		-40000000.00
1	4882828.70	-35117171.30	.9524	4650313.26
2	5075527.48	-30041643.82	.9070	4603653.46
3	5288404.34	-24753239.47 -19229667.28 -13446302.09	.8638	4568323.13
4	5523572.19	-19229667.28	.8227	4544257.34
5	5783365.19	-13446302.09	.7835	4531418.98
		-7715042.34		
		-1327631.34		
		3180019.78	.6768	
9	4233380.00 8607849.67			2728875.61 5284472.96
10	860/849.6/	16021249.45	.0133	52844/2.96
THE NET	PRESENT VALUE	(\$): 427784	442.19	
PRESENT	VALUE RATIO:	1.069		
THE NET	RETURN RATE:	10.695	8	
THE AVE	RAGE RATE OF RE	TURN: 14.005	¥	
THE PAY	BACK PERIOD IS	BETWEEN: 7	AND 8 YEARS	

Table 9-16 Input Data and Computer Output for the Discount Cash Flow Rate of Retern (DCFRR) the the Economic Project Evaluation

DATA92.DAT			
11			
-40000000.0			
5075527.48			
5288404.34			
5523572.19			
5783365.19 5731259.75			
6387411.00			
4507651.13			
4233380.00			
8607849.67			
		NT CASH FLOW RATE OF RETURN CALCULAT	
*****		NT CASH FLOW RATE OF RETURN CALCULAT	

*****	******		
*****	******	***************	
	**************************************	CASH FLOWS INCLUDING YEAR 0	*****
	**************************************	CASH FLOWS INCLUDING YEAR 0 CASH FLOW (\$) -40000000.00	*****
	**************************************	**************************************	*****
	**************************************	CASH FLOWS INCLUDING YEAR 0 CASH FLOW (\$) -40000000.00	*****
	**************************************	**************************************	*****
	**************************************	**************************************	*****
	**************************************	**************************************	*****
	**************************************	**************************************	*****
	**************************************	**************************************	*****

THE DISCOUNT CASH FLOW RATE OF RETURN (%): 6.347

(text continued from page 754)

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PROGRAM PROG91

~	*********
С	······
с	* THIS PROGRAM CALCULATES THE YEARLY CASH FLOWS, THE CUMULATIVE *
С	* CASH FLOWS, DISCOUNT FACTORS AT A GIVEN RATE AND PRESENT *
с	* VALUES. THE PROGRAM FURTHER CALCULATES THE NET PRESENT VALUE, *
č	* PRESENT VALUE RATIO, THE AVERAGE RATE OF RETURN, THE NET *
С	* RETURN RATE, AND THE PAYBACK PERIOD. *
С	***************************************
с	ARR = AVERAGE RATE OF RETURN YCF = YEARLY CASH FLOW DF = DISCOUNT FACTOR
С	YCF = YEARLY CASH FLOW
с	DF = DISCOUNT FACTOR
с	CUCF = CUMULATIVE CASH FLOW
č	NDV = NET DDESENT VALUE
č	NOT * NIMBED OF DOLLECT LIFE FYCHIDING YEAD O
	NOCT - NOADER OF FRONECT LIFE EACEODING TEAR O
С	NRR = NET RETURN RATE
С	DIF = DIFUSION FACTOR CUCF = CUMULATIVE CASH FLOW NPV = NET PRESENT VALUE NOCF = NUMBER OF PROJECT LIFE EXCLUDING YEAR 0 NRR = NET RETURN RATE PV = PRESENT VALUE
с	PVR = PRESENT VALUE RATIO
с	TIME = PAYBACK PERIOD
С	***************************************
	REAL*8 YCF, DF, PV, NPV, CUCF
	EXTERNAL XNPV REAL NRR
	INTEGER TIME, TIME1
с	DIMENSION YCF(0:100), DF(0:100), PV(0:100), CUCF(0:100)
	OPEN (UNIT = 3, FILE = 'DATA91.DAT', STATUS ='OLD', ERR=18) OPEN (UNIT = 1, FILE = 'PRN')
с	READ THE NUMBER OF YEARLY CASH FLOWS EXCLUDING YEAR 0: NOCF
	READ (3, *, ERR=19) NOCF
C C	READ THE ANNUAL DISCOUNT RATE IN PERCENTAGE: DISC READ THE YEARLY CASH FLOW: YCF
	READ (3, *, ERR=19) DISC READ (3, *, ERR=19) (YCF(K), K = 0, NOCF)
	GO TO 10
18 21	WRITE (*, 21) Format (6x,'data file does not exist') Go to 999
19 23	WRITE (*, 23) Format(6x,'error message in the data value') Go to 999

```
10
         WRITE (1, 100)
FORMAT (//,25X,'NET PRESENT VALUE CALCULATION',/1H ,78(1H*))
 100
         NOCF1 = NOCF + 1
         WRITE (1, 110) NOCF1
         FORMAT(/, 20X, 14, 3X, 'YEARLY CASH FLOWS INCLUDING YEAR 0')
 110
         WRITE (1, 120) DISC
         FORMAT (/,20X,F8.2,3X,'PERCENTAGE ANNUAL DISCOUNT RATE',/,
 120
        1H ,78(1H-))
         WRITE (1, 130)
FORMAT (/,7X,'YEAR',5X,'CASH FLOWS ($)',4X,'CUMULATIVE',\)
 130
         WRITE (1, 140)
FORMAT (10X, 'DISCOUNT', 8X, 'PRESENT')
 140
         WRITE (1, 150)
        FORMAT (34X, 'CASH FLOWS ($)', 6X, 'FACTOR', 10X, 'VALUE ($)',
 150
                   /1H ,78(1H-))
         CALCULATE THE CUMULATIVE CASH FLOW
с
         CUCF(0) = YCF(0)
         DO I = 1, NOCF
         CUCF(I) = CUCF(I-1) + YCF(I)
         END DO
         R1 = DISC/100.0
         PV(0) = YCF(0) * 1.0
         DO 15 I1 = 0, 0
         IF (I1 .EQ. 0) THEN R1 = 0.0
         DF(I1) = 1.0/((1.0+R1)**I1)
         PV(I1) = YCF(I1) * DF(I1)
         ELSE
         ENDIF
         CONTINUE
15
         R1 = DISC/100.0
         DO 20 I1 = 1, NOCF
DF(I1) = 1.0/((1.0+R1)**I1)
PV(I1) = YCF(I1)*DF(I1)
         CONTINUE
  20
         DO K = 0, NOCF
WRITE (1, 160) K, YCF(K), CUCF(K), DF(K), PV(K)
FORMAT (5X,I3,5X,F14.2,5X,F14.2,5X,F9.4,5X,F14.2)
160
         END DO
         WRITE (1, 170)
170
         FORMAT (78(1H-))
```

```
NPV = 0.0
         DO J = 0, NOCF
NPV = NPV + PV(J)
         END DO
         WRITE (1, 180) NPV
         FORMAT (1HO, 6X, 'THE NET PRESENT VALUE ($):', 3X, T40, F14.2)
 180
         CALCULATE THE PRESENT VALUE RATIO, PVR
с
č
         PVR = PRESENT VALUE OF ALL POSITIVE CASH FLOWS/
PRESENT VALUE OF ALL NEGATIVE CASH FLOWS
         SUM1 = 0.0
         DO K = 1, NOCF
SUM1 = SUM1 + PV(K)
         END DO
         PVR = SUM1 /ABS(YCF(0))
         WRITE (1, 190) PVR
         FORMAT (1H0, 6X, 'PRESENT VALUE RATIO:', 3X, T40, F8.3)
 190
с
         CALCULATE THE NET RETURN RATE (NRR) i.e. THE NET PRESENT VALUE
         UNIDED BY THE PRODUCT OF INITIAL INVESTMENT AND THE PROJECT LIFE (EXCLUDING YEAR 0).
c
c
         NRR = ABS(NPV / (ABS(PV(0)) * NOCF)) * 100
         WRITE (1, 200) NRR
         FORMAT (1H0, 6X, 'THE NET RETURN RATE:', 3X, T40, F8.2, 3X, '%')
 200
С
         CALCULATE THE AVERAGE RATE OF RETURN (ARR) i.e. THE AVERAGE
         CASH FLOW DURING THE LIFE OF THE PROJECT (EXCLUDING YEAR 0).
с
        SUM = 0.0
DO I = 1, NOCF
SUM = SUM + YCF(I)
         END DO
         AVSUM = SUM/(NOCF)
ARR = (AVSUM/ABS(YCF(0)))*100.0
         WRITE (1, 210) ARR
 210
         FORMAT (1H0,6X, 'THE AVERAGE RATE OF RETURN: ', 3X, T40, F8.2,
                 3x, (%()
с
         CALCULATE THE PAYBACK TIME FROM THE CUMULATIVE CASH FLOW, TIME: YEARS
         DO I = 0, NOCF
         IF ( CUCF(I+1) .GT. CUCF(I) .AND. CUCF(I+1) .GE. 0.0) THEN
```

```
TEMP = CUCF(I+1)
CUCF(I+1) = CUCF(I)
CUCF(I) = TEMP
TIME = I
TIME1 = TIME + 1
GO TO 40
ELSE
ENDIF
END DO
40 WRITE (1, 220) TIME, TIME1
220 FORMAT (1H0,6X,'THE PAYBACK PERIOD IS BETWEEN:',
* 3X,T45,I2,2X,'AND',2X,I2,3X,'YEARS')
C FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
WRITE (1, *) CHAR(12)
CLOSE (3, STATUS = 'KEEP')
CLOSE (1)
999 STOP
END
```

PROGRAM PROG92

с с с	* THIS PROGRAM CALCULATES THE DISCOUNT CASH FLOW RATE OF * RETURN FROM A SET OF YEARLY CASH FLOWS *
00000	YCF = YEARLY CASH FLOW DCFRR = DISCOUNT CASH FLOW RATE OF RETURN NOCF = THE NUMBER OF YEARLY CASH FLOW INCLUDING YEAR 0
	EXTERNAL DCFRR
	DIMENSION YCF (1:100)
	OPEN (UNIT = 3, FILE = 'DATA92.DAT', STATUS = 'OLD', ERR = 18) OPEN (UNIT = 1, FILE = 'PRN')
с	READ THE NUMBER OF YEARLY CASH FLOWS INCLUDING YEAR 0: NOCF
	READ (3, *, ERR=19) NOCF
с	READ THE YEARLY CASH FLOWS: YCF
	READ (3, \star , ERR=19) (YCF(K), K = 1, NOCF)
	GO TO 10
18 21	WRITE (*, 21) Format (6x, 'data file does not exist') Go to 999
19 23	WRITE (*, 23) Format (6x, 'error message in the data value') Go to 999
10 100 *	<pre>WRITE (1, 100) FORMAT (///,20X,'DISCOUNT CASH FLOW RATE OF RETURN CALCULATION',</pre>
110	WRITE (1, 110) NOCF FORMAT (/,1H0,15X,I4,3X,'YEARLY CASH FLOWS INCLUDING YEAR 0')
120	WRITE (1, 120) FORMAT (//, 17X,'YEAR', 12X, 'CASH FLOW (\$)', /1H ,78(1H*))
	DO 20 K = 1, NOCF
130 20	L = K - 1 WRITE (1, 130) L, YCF(K) FORMAT (15X, I3, 11X, F14.2) CONTINUE WRITE (1, 140)

```
140
        FORMAT (78(1H-))
        DCF = DCFRR (YCF, NOCF)
        WRITE (1, 150) DCF
FORMAT (/,1H0,6X,'THE DISCOUNT CASH FLOW RATE OF RETURN (%):',
3X, T50, F8.3)
 150
С
        FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.
        WRITE (1, *) CHAR(12)
        CLOSE (3, STATUS = 'KEEP')
        CLOSE (1)
 999
        STOP
        END
С
С
        THIS PROGRAM CALCULATES THE DISCOUNT CASH FLOW RETURN OF A SET
С
c
        OF YEARLY CASH FLOWS
        FUNCTION DCFRR (CF, IYRS)
        DIMENSION CF (1:100)
        XIF = 1.
        M = 1
        PV = CF(IYRS)
10
        IF (M .GT. 100) THEN
GO TO 20
        ELSE
        DO 30 J = 1, IYRS - 1
PV = PV*XIF + CF(IYRS-J)
        CONTINUE
30
        ENDIF
        DPV = (IYRS-1)*CF(IYRS)
        DO 40 J = 1, IYRS - 2
DPV = DPV*XIF + (IYRS-J-1)*CF(IYRS-J)
        CONTINUE
40
        P = PV/DPV
        IF ((ABS(P)-1.E-4) .GT. 0.0) THEN
        GO TO 50
        ELSE
       X1 = 1/XIF - 1.
DCFRR = 100 * X1
        GO TO 60
       ENDIF
50
       XIF = XIF - P
       M = M + 1
       GO TO 10
       WRITE (1, 160)
FORMAT (//,1H0,3X,'DCFRR CONVERGENCE OF NEXT CASH FLOW DOES ,
OCCUR AFTER 100 TRIES',/)
20
160
       RETURN
60
        END
```

PROGRAM PROG93

	<pre>************************************</pre>
С	REAL*8, B, E, S1, BT, X, CF, CUCF, DF, PV, NPV
	DIMENSION X(1:100), CF(0:100), CUCF(0:100), DF(0:100), PV(0:100)
	EXTERNAL FUN
	INTEGER A, FL, N, TIME, TIME1 Real NRR
	OPEN (UNIT = 3, FILE = 'DATA93.DAT', STATUS = 'OLD', ERR = 18) OPEN (UNIT = 1, FILE = 'PRN')
	READ (3, *, ERR=19)(X(I),I=1,9) READ (3, *, ERR=19)R
100 *	WRITE (1, 100) FORMAT (///, 22X,'ECONOMIC EVALUATION OF A PROJECT', /1H ,78(1H*))
	I = 1
110	WRITE (1, 110) I, X(I) FORMAT (1H0, 6X, I2, 6X,'ANNUAL REVENUE \$:',T55, F14.2)

	I = I+1
120	WRITE (1, 120) I, X(I) FORMAT (1H0, 6X, I2, 6X, 'ANNUAL OPERATING COST \$:',T55, F14.2)
	I = I + 1
130	WRITE (1, 130) I, X(I) FORMAT (1H0, 6X, I2, 6X, 'DEPRECIATION BASE \$:',T55, F14.2)
	I = I + 1
140	WRITE (1, 140) I, X(I) FORMAT (1H0, 6X, I2, 6X, 'PROJECT LIFE, YEARS:',T55, F3.0)
	$I \neq I + 1$
150	WRITE (1, 150) I, X(I) Format (1h0, 6X, I2, 6X, 'INITIAL LOAN \$:',T55, F14.2)
	I = I + 1
160	WRITE (1, 160) I, X(I) FORMAT (1H0,6X,I2,6X,'NUMBER OF PAYMENTS PER YEAR:',T55,F4.0)
	I = I + 1
170	WRITE (1, 170) I, X(I) FORMAT (1H0, 6X, I2, 6X, 'ANNUAL INTEREST, %:',T55, F4.0)
	I = I + 1
180	WRITE (1, 180) I, X(I) FORMAT (1H0, 6X, I2, 6X, 'INVESTMENT CREDIT, %:',T55, F4.0)
	I = I + 1
190	WRITE (1, 190) I, X(I) Format (1H0, 6X, I2, 6X, 'TAX RATE, %:',T55, F4.0)
	I = I + 1
200	WRITE (1, 200) I, R FORMAT (1H0, 6X, I2, 6X, 'DISCOUNT RATE %:',T55, F6.2)
210	WRITE (1, 210) FORMAT (78(1H-))
	GO TO 20
18 220	WRITE (*, 220) FORMAT (6X, 'DATA FILE DOES NOT EXIST')
	GO TO 999
19 230	WRITE (*, 230) Format (6x, 'error message in the data value')
	GO TO 999

20 22	A = 0 FL = 0 IF (FL .EQ. 1) THEN GO TO 60 ELSE
	Q7 = X(3) $Q5 = X(5)$ $Q2 = X(7)$ $X(7) = X(7)/(100.0*X(6))$ $Q1 = X(8)$ $X(8) = (X(8)*X(3))/100.0$ $P3 = (X(5)*X(7))/(1.0-(1.0+X(7))**(-X(4)*X(6)))$ ENDIF
21 25	$\begin{array}{l} \lambda = \lambda + 1 \\ D = (2.0*X(3))/X(4) \\ X(3) = X(3) - D \\ I = X(6) \\ B = B + (X(5)*X(7)) \\ VAL1 = Q5 + (X(5)*X(7)) - P3 \end{array}$
	IF (VAL1 .GT. 0.0) THEN GO TO 30 ELSE
240	WRITE (1, 240) Format (1H0, 6X, 'ERROR IN THE DATA VALUE')
250 *	WRITE (1, 250) FORMAT (1H0,6X,'I.E. THE TOTAL LOAN PLUS THE INTEREST PER',/1H , 6X,'PERIOD IS LESS THAN THE PAYMENTS') ENDIF
	GO TO 999
30	X(5) = X(5) + (X(5) * X(7)) - P3
	I = I - 1
	IF (I .LT. 0 .OR. I .GT. 0) THEN GO TO 25 ELSE
260	WRITE (1, 260) A Format (/, 1H1, 32X, 'YEAR', 2X, I2, /1H , 78(1H*))
270	WRITE (1, 270) X(1) FORMAT (1H0, 6X, 'REVENUE \$:', T55, F14.2)
280	WRITE (1, 280) X(2) FORMAT (1H0, 6X, 'OPERATING COST \$:', T55, F14.2)
290	WRITE (1, 290) D Format (1H0, 6X, 'DEPRECIATION \$:', T55, F14.2)
300	WRITE (1, 300) B Format (1H0, 6X, 'INTEREST \$:', T55, F14.2)

```
ENDIF
        E = X(1) - X(2) - D - B
        X1 = E
IF (X1 .LT. 0.0) THEN
GO TO 40
        ELSE
        X1 = E - PO
        ENDIF
        IF (X1 .LT. 0.0) THEN
GO TO 50
        ELSE
        E = X1
        PO = 0.0
        S1 = (E*X(9))/100.0
X1 = S1 - X(8)
        ENDIF
        IF (X1 .EQ. 0.0 .OR. X1 .GT. 0.0) THEN GO TO 55
        ELSE
        X(8) = X(8) - S1
        S1 = 0.0
        ENDIF
        WRITE (1, 310) E
FORMAT (1H0, 6X, 'PRE-TAX INCOME $:', T55, F14.2)
70
310
        WRITE (1, 320) X(9), S1
        FORMAT (1H0, 6X, 'TAX AT', F4.0,'%, $:', T55, F14.2)
320
        BT = E - S1
        WRITE (1, 330) BT
        FORMAT (1H0, 6X, 'AFTER TAX INCOME $:', T55, F14.2)
330
        CF(A) = BT + D
        WRITE (1, 340) CF(A)
FORMAT (1H0, 6X, 'CASH FLOW $:', T55, F14.2)
340
        WRITE (1, 350)
FORMAT ( 78(1H*))
350
        B = 0.0
        IF ((A-X(4)) .LT. 0.0) THEN
        GO TO 21
        ELSE
        WRITE (1, 360) X(3)
FORMAT (1H0, 6X, 'SALVAGE VALUE $:', T55, F14.2)
360
        ENDIF
        GO TO 60
        PO = PO - X1
  40
```

```
S1 = 0.0
```

50	GO TO 70 PO = -X1 X1 = 0.0 GO TO 40
55	S1 = X1 X(8) = 0.0 GO TO 70 FL = 1
	GO TO 22
с	CALCULATE THE NET PRESENT VALUE OF INVESTMENT
60 370 *	<pre>wRITE (1, 370) FORMAT (1H1,30X,'ECONOMIC PROJECT EVALUATION',/1H ,20X,</pre>
	NOCF = $X(4) + 1$ N = $X(4)$
380	WRITE (1, 380) NOCF FORMAT (/,1H0, 22X,I3,3X,'YEARLY CASH FLOWS INCLUDING YEAR 0')
390 *	<pre>WRITE (1, 390) R FORMAT (/,1H0,20X,F8.2,3X,'PERCENTAGE ANNUAL DISCOUNT RATE',/, 1H , 78(1H-))</pre>
400 *	WRITE (1, 400) FORMAT (/, 1H0, 15X,'YEARLY CASH FLOWS (\$)',/1H ,78(1H-),/1H0, 7X, 'YEAR', 5X, 'CASH FLOWS (\$)', 3X, 'CUMULATIVE',\)
410	WRITE (1, 410) Format (10x, 'DISCOUNT', 8x, 'PRESENT')
420 *	WRITE (1, 420) FORMAT (34X, 'CASH FLOWS (\$)', 6X, 'FACTOR', 10X, 'VALUE (\$)', /1H, 78(1H-))
с	CALCULATIVE THE CUMULATIVE CASH FLOW
	CUCF(0) = -Q7
	DO I = 1, N-1 CUCF(I) = CUCF(I-1) + CF(I) END DO
	CF(N) = CF(N) + X(3)
	CUCF(N) = CUCF(N-1) + CF(N)
	R1 = R/100.0 CF(0) = -Q7 PV(0) = CF(0)*1.0

C CALCULATE THE PRESENT VALUE FOR YEAR 0

DO 75 I1 = 0, 0 IF (I1 .EQ. 0) THEN R1 = 0.0DF (I1) = 1.0/((1.0+R1)**I1) PV(I1) = CF(I1) * DF(I1)ELSE ENDIF 75 CONTINUE CALCULATE THE PRESENT VALUE FOR SUBSEQUENT YEARS с R1 = R/100DO 80 I1 = 1, N-1 с I2 = I1 - 1DF (I1)= 1.0/((1.0+R1)**I1) PV(I1) = CF(I1)*DF(I1) 80 CONTINUE C C CALCULATE THE PRESENT VALUE FOR THE FINAL YEAR PLUS THE SALVAGE VALUE DF(N) = 1/((1+R1)**N)PV(N) = CF(N)* DF(N)DO 90 K=0, N PRINT THE VALUES OF THE YEARLY CASH FLOWS, CUMULATIVE CASH FLOWS c c DISCOUNT FACTORS AND THE PRESENT VALUES WRITE (1, 430) K, CF(K), CUCF(K), DF(K), PV(K) FORMAT (5X, I3, 5X, F14.2, 5X, F14.2, 5X, F9.4, 5X, F14.2) 430 CONTINUE 90 WRITE (1, 440) 440 FORMAT (78(1H-)) с CALCULATE THE NET PRESENT VALUE NPV = 0.0DO J = 1, NOCF NPV = NPV + PV(J) END DO WRITE (1, 450) NPV FORMAT (1H0, 6X, 'THE NET PRESENT VALUE (\$):',3X,T40,F14.2) 450 CALCULATE THE PRESENT VALUE RATIO, PVR С PVR = PRESENT VALUE OF ALL POSITIVE CASH FLOWS/ = PRESENT VALUE OF ALL NEGATIVE CASH FLOWS c c

```
SUM = 0.0
         DO K = 1, N
SUM = SUM + PV(K)
         END DO
         PVR = SUM/ABS(CF(0))
         WRITE (1, 460) PVR
 460
         FORMAT (1H0, 6X, 'PRESENT VALUE RATIO:', 3X, T40, F8.3)
         CALCULATE THE NET RETURN RATE (NRR) i.e. THE NET PRESENT VALUE
С
         DIVIDED BY THE PRODUCT OF INITIAL INVESTMENT AND THE PROJECT LIFE (EXCLUDING YEAR 0).
С
С
         NRR = ABS (NPV/(ABS(CF(0))*N))*100
         WRITE (1, 470) NRR
 470
         FORMAT (1H0,6X,'THE NET RETURN RATE:',3X,T40,F8.3,3X,'%')
         CALCULATE THE AVERAGE RATE OF RETURN (ARR) i.e. THE AVERAGE CASH FLOW DURING THE LIFE OF THE PROJECT (EXCLUDING YEAR 0).
С
С
         SUM1 = 0.0
         DO I = 1, N
          SUM1 = SUM1 + CF(I)
         END DO
         AVSUM = SUM1/N
         ARR = (AVSUM/ABS(CF(0)))*100.0
         WRITE (1, 480) ARR
480
         FORMAT (1H0, 6X, 'THE AVERAGE RATE OF RETURN:', 3X, T40, F8.3, 3X, '%')
c
c
         CALCULATE THE PAYBACK TIME FROM THE CUMULATIVE CASH FLOW
         TIME: Years
         DO I = 0, N
         IF (CUCF(I+1) .GT. CUCF(I) .AND. CUCF(I+1) .GE. 0.0) THEN
         TEMP = CUCF (I+1)
CUCF (I+1) = CUCF (I)
CUCF (I) = TEMP
TIME = I
         TIME1 = TIME +1
         GO TO 44
         ELSE
         ENDIF
         END DO
         WRITE (1, 490) TIME, TIME1
FORMAT (1H0,6X,'THE PAYBACK PERIOD IS BETWEEN:',
 44
 490
                  3X,T45,I2,2X,'AND',2X,I2,2X,'YEARS')
```

C FORM FEED THE PRINTING PAPER TO TOP OF THE NEXT PAGE.

```
WRITE (1, *) CHAR(12)
CLOSE (3, STATUS = 'KEEP')
CLOSE (1)
STOP
END
```

999

CHAPTER 10

The International System of Units (SI) and Conversion Tables

The International System of Units, or the SI units, are derived from Systeme International d'Unites and are evolved from the mass-lengthtime-temperature system. This is derived from the seven base units of length, mass, time, temperature, amount of substance, electric current, and luminous intensity. Also, two supplementary units of plane angle and solid angle are required to describe the physical and chemical properties. Table 10-1 lists the supplementary units and their symbols.

All other units in SI are derived from the nine base and supplementary units. Table 10-2 gives approved derived units with names, formulas, and their symbols. Lists of derived SI units that do not have approved names but are widely used in chemical engineering are given in Table 10-3. The wide range in magnitude of the units used in normal practice requires the use of different sized units, rather than a powers-of-ten numerical multiplier. SI provides prefixes for decimal multiples and submultiples of the base supplementary and approved derived units. Table 10-4 shows these prefixes by name, symbol, and magnitude. In addition to learning the new units and symbols, conversion is required when numerical values are reported in imperial or the c.g.s units. The conversion of units is mainly handled through the use of conversion tables as listed in Table 10-5.

Туре	Quantity and Symbol	Unit	Unit Symbol	
base	Length, L	meter m		
base	mass, m	kilogram	kg	
base	time, t	second	S	
base	temperature, T	Kelvin	K	
base	amount of substance, n	mole	mol	
base	electric current, I	ampere	А	
base	luminous intensity, I	candela	cd	
Supplementary	plane angle, θ	radian	rad	
Supplementary	solid angle, ω	steradian	sr	

Table 10-1SI and Supplementary Units

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Quantity and symbol	Unit	Unit symbol	Formula
frequency, f	hertz	Hz	s ⁻¹
force, F	newton	N	kg.m/s ²
pressure, stress p, τ	pascal	Ра	N/m², kg/m.s²
energy, work, quantity of heat, E, W, Q	joule	J	N/m, kg.m²/s²
power, radiant flux, P	watt	W	J/s, kg.m ² /s ³
quantity of electricity, electric charge, Q	coulomb	С	A.s
electric potential, potential difference, electromotive force, E	volt	V	W/A, kg/A.s ³
capacitance, C	farad	F	C/V, A ² .s ⁴ /kg.m ²
electric resistance, R	ohm	Ω	V/A, m ² .kg/A ² .s ³
conductance, G	siemens	S	A/V, A ² .s ³ /kg.m ²
magnetic flux, ø	weber	Wb	V.s, m ² .kg/A.s ²
magnetic flux density, B	tesla	Т	Wb/m ² , kg/A.s ²
inductance, L	henry	Н	Wb/A, m ² .kg/A.s
luminous flux, F	lumen	lm	cd.sr
illuminance, B	lux	lx	cd.sr/m ²
radioactivity, λ	becquerel	Bq	s ⁻¹
absorbed dose, D	gray	Gy	J/kg

Table 10-2SI Derived Units with Special Names

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Table 10-3 Some Derived SI Units

Property	Common Symbol	SI units formula
thermal conductivity	k	W/m.K
heat transfer rate	q	W
heat transfer coefficient	h	W/m ² .K
specific enthalpy	Н	J/kg
heat capacity	C _p	J/kg.K
Newtonian viscosity	μ	Pa.s, kg/m.s
kinematic viscosity	ν	m²/s
thermal diffusivity	α	m ² s
volumetric flow rate	q	m³/s
power	Р	W
enthropy	S	J/K
density	ρ	kg/m ³
mass velocity	G	kg/s.m ²
concentration	с	kg/m ³
concentration	с	mol/m ³
velocity	u	m/s
acceleration	a,g	m/s ²
mass transfer coefficient	k _c	m/s
mass diffusivity	D	m²/s
moment	Ι	N.m
thermal flux	q/A	W/m ²
mass flux (molar)	N _a /A	mol/m ² .s
momentum flux	τ_y/A	Ра
mass transfer rate	N _a	mol/s
mass flow rate	w	kg/s
surface tension	γ	N/m
gas constant	R	Pa.m ³ /mol.K, J/mol.K

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Name	Symbol	Magnitude
atto	a	10-18
femto	f	10-15
pico	р	10 ⁻¹²
nano	n	10-9
micro	μ	10-6
milli	m	10-3
centi	с	10-2
deci	d	10-1
deka	da	101
hecto	h	102
kilo	k	103
mega	M	106
giga	G	109
tera	Т	1012
peta	Р	1015
exa	E	1018

Table 10-4Magnitude Prefixes for SI Units

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Table 10-5 Conversion of Units

			Length			
in	ft	mm	cm	m	km	mile
1	0.083333	25.4	2.54	0.0254		
12	1	304.8	30.48	0.3048	3.048×10^{-4}	1.894 × 10-
0.03937	3.2808×10^{-3}	1	0.1	10-3		
0.3937	0.032808	10	1	0.01		
39.3701	3.28084	1000	100	1	10-3	6.214 × 10⁻
39370.1	3280.84	106	100000	1000	1	0.621371
63360	5280	1.60934×10^{6}	160934	1609.34	1.60934	1

		111.04	•		
in ²	ft2	mm ²	cm ²	m ²	km²
1	6.944 × 10 ⁻³	645.16	6.4516	6.452×10^{-4}	
144	1	92903	929	0.0929	
1.55 × 10 ⁻³	1.076×10^{-5}	1	0.01	10-6	
0.155	1.076×10^{-3}	100	1	10-4	
1550	10.764	106	10000	1	
	1.0764×10^{7}			106	1

Area

(table continued on next page)

Table 10-5	
(continued)	

			Volume			
in ³	ft3	mm ³	ml	l	m ³	US gal
I	5.787 × 10 ⁻⁴	16387	16.39	0.01639	1.639 × 10 ⁻⁵	4.329×10^{-3}
1728	1	2.832×10^{7}	2.832×10^{4}	28.32	0.02832	7.4805
6.1024 × 10 ⁻⁵	3.531 × 10 ⁻⁸	1	1 × 10 ⁻³	10-6	10-9	2.642×10^{-7}
0.061026	3.532 × 10 ⁻⁵	1000	1	10-3	10-6	2.642×10^{-4}
61.026	0.03532	1 × 10 ⁶	1000	1	1×10^{-3}	0.2642
61024	35.31	109	1×10^{6}	1000	1	264.2
231.0	0.1337	3.785×10^{6}	3785	3.785	3.785×10^{-3}	1

1 US barrel = 42 US gal = 34.97 UK gal 1 US gal = 0.832674 UK gal

Mass					
lb	g	kg	t (tonne)	US ton (short ton)	
1	453.592	0.453592	4.5359 × 10 ⁻⁴	5.0×10^{-4}	
2.2046×10^{-3}	1	10-3		<u> </u>	
2.20462	103	1	10-3	1.1023×10^{-3}	
2204.62	106	1000	1	1.10231	
2000	907185	907.185	0.907185	1	

(table continued on next page)

Table 10-5 (continued)

Density					
lb/in ³	lb/ft ³	g/cm ³	kg/m³		
1	1728	27.6799	27679.9		
5.787 × 10 ⁻⁴	1	0.0160185	16.0185		
0.036127	62.428	1	1000		
3.6046 × 10 ⁻⁵	0.062428	10-3	1		

	Velocity									
ft/s	ft/min	mm/s	cm/s	m/s	km/s	km/h				
1	60	304.8	30.48	0.3048	3.048×10^{-4}	1.09728				
0.016667	1	5.08	0.508	5.08×10^{-3}	5.08 × 10 ⁻⁶	0.018288				
3.281 × 10 ⁻³	0.19685	1	0.1	10-3	10-6	3.6×10^{-3}				
0.032808	1.9685	10	1	0.01	10-5	0.036				
3.28084	196.85	1000	100	1	10-3	3.6				
3280.84	196850	106	100000	103	1	3600				
0.911344	54.6806	277.778	27.7778	0.277778	2.778×10^{-4}	1				

Table 10-5 (continued)

	Temperature					
Kelvin, K	$T = \theta + 273.15 = \frac{5}{9}(t + 459.67) = \frac{5}{9}(r)$					
degree Celsius, °C	$t = T - 273.15 = \frac{5}{9}(t - 32) = \frac{5}{9}(r - 491.67)$					
degree Fahrenheit, °F	$t = \frac{9}{5}(T) - 459.67 = \frac{9}{5}(\theta) + 32 = r - 459.67$					
degree Rankine, °R	$\mathbf{r} = \frac{9}{5}(\mathbf{T}) = \frac{9}{5}(\mathbf{\theta}) + 491.67 = \mathbf{t} + 459.67$					

where T = Kelvin, $\theta = {}^{\circ}C$, $t = {}^{\circ}F$, $r = {}^{\circ}R$

lb/s	lb/min	lb/h	g/s	kg/s	kg/h	tonne/d
1	60	3600	453.6	0.4536	1633	39.19
1.667×10^{-2}	1	60	7.560	7.56×10^{-3}	27.216	0.653
2.778×10^{-4}	0.01667	1	0.1260	1.26×10^{-4}	0.4536	0.01089
2.205×10^{-3}	0.1323	7.937	1	1 × 10 ⁻³	3.60	0.08640
2.205	132.3	7937	1000	1	3600	86.40
6.124 × 10 ⁻⁴	0.03674	2.205	0.2778	2.778 × 10 ⁻⁴	1	0.02400
0.02551	1.531	91.86	11.57	0.01157	41.67	1

Mass Flow Rate (mass/time)

Table 10-5 (continued)

		Volur	netric Flow Rat	te (volume/tin	ne)		
ft³/s	ft³/min	US gal/min	ml/s	l/s	l/min	m³/s	m³/h
1	60	448.83104	2.832×10^{4}	28.32	1699	0.02832	101.9
0.01667	1	7.480717	471.9	0.4719	28.31	4.719×10^{-4}	1.699
2.228×10^{-3}	0.133644	1	63.0917	0.063092	3.78534	6.3092 × 10 ⁻⁵	0.22715
3.532×10^{-5}	2.119×10^{-3}	0.158503	1	1×10^{-3}	0.06	1 × 10 ⁻⁶	3.6×10^{-3}
0.03531	2.119	15.85026	1000	1	60	1×10^{-3}	3.6
5.886×10^{-4}	0.03531	0.264172	16.67	0.01667	1	1.667×10^{-5}	0.0600
35.31	2119	1.585×10^{4}	1 × 10 ⁶	1000	6×10^{4}	1	3600
9.81 × 10 ⁻³	0.5886	4.40286	277.8	0.2778	16.67	2.778×10^{-4}	1

1 US gal/min = 0.832674 U.K. gal/min

1 UK gal/min = 1.20095 US gal/min

	Dynamic Viscosity									
pdls/ft2	lb/ft.h	сP	Р	kg/m.h	Ns/m ²					
1	3600	1488	14.88	5358	1.488					
2.776 × 10 ⁻⁴	1	0.4134	4.134×10^{-3}	1.488	4.134×10^{-4}					
6.720 × 10 ⁻⁴	2.419	1	0.01	3.6	10-3					
0.0672	241.9	100	1	360	0.100					
1.866 × 10 ⁻⁴	0.6720	0.2778	2.778×10^{-3}	1	2.788×10^{-4}					
0.6720	2419	1000	10	3600	1					

Table 10-5 (continued)

Force									
poundal pdl	pound-force lbf	kilogram-force kgf	newton N	KN					
1	0.0311	0.0141	0.1383	1.383×10^{-4}					
32.174	1	0.4536	4.448	4.448×10^{-3}					
70.93	2.2046	1	9.807	9.807×10^{-3}					
7.233	0.2248	0.1020	1	10-3					
7233	224.8	102.0	1000	1					

Foot pound-force ft lbf	British thermal unit Btu	calorie cal	horsepower hour hph	kilowatt hour kWh	Joule J	kilojoule kJ
1	1.285×10^{-3}	0.3238	5.051×10^{-7}	3.766×10^{-7}	1.3558	1.356×10^{-3}
778.17	1	252	3.930×10^{-4}	2.931 × 10 ⁻⁴	1055.1	1.0551
3.0880	3.968×10^{-3}	1	1.560×10^{-6}	1.163 × 10-6	4.1868	4.187×10^{-3}
1.98×10^{6}	2544.4	641186	1	0.7457	2.6845×10^{6}	2684.5
2.65522×10^{6}	3412.1	859845	1.3410	1	3.600×10^{6}	3600
0.7376	9.478 × 10 ⁻⁴	0.2388	3.725×10^{-7}	2.778 × 10 ⁻⁷	1	10-3
737.56	0.9478	238.85	3.725×10^{-4}	2.778×10^{-4}	1000	1

Energy, Work, Heat

Table 10-5 (continued)

	Pressure										
lb _f /in ²	lb _f /ft²	in H ₂ O	mm Hg	atm	Pascal, Pa N/m²	kN/m²	bar				
1	144	27.68	51.71	0.06805	6895	6.895	0.06895				
6.944×10^{-3}	1	0.1922	0.3591	4.726×10^{-4}	47.88	0.04788	4.788 × 10 ⁴				
0.03613	5.202	1	1.868	2.458×10^{-3}	249.1	0.2491	2.491×10^{-3}				
0.01934	2.785	0.5352	1	1.316×10^{-3}	133.3	0.1333	1.333×10^{-3}				
14.70	2116	406.8	760.0	1	1.013×10^{5}	101.3	1.013				
1.450×10^{-4}	0.02089	4.015×10^{-3}	7.501×10^{-3}	9.869 × 10 ⁻⁶	1	1×10^{-3}	1 × 10 ⁻⁵				
0.1450	20.89	4.015	7.501	9.869×10^{-3}	1000	1	0.0100				
14.50	2089	401.5	750.1	0.9869	1×10^{5}	100	1				

ft lb _f /lb°F	Btu/lb°F	kcal/kg.K	kgf m/kgK	J/kg.K	kJ/kg.K
1	1.28507×10^{-3}	1.28507×10^{-3}	0.54864	5.38032	5.380×10^{-3}
778.169	1	1	426.935	4186.8	4.1868
778.169	1	1	426.935	4186.8	4.1868
1.82269	2.34228×10^{-3}	2.34228×10^{-3}	1	9.80665	9.807 × 10 ⁻³
0.185863	0.2388×10^{-3}	0.2388 × 10 ⁻³	0.10197	1	10-3
185.863	0.2388	0.2388	101.97	1000	1

Specific Heat Capacity (heat/mass × degree temperature)

Table 10-5 (continued)

Power, Heat Flow Rate									
ft lbf/s	Btu/h	hp	kcal/h	W	kW				
1	4.626	1.818×10^{-3}	1.1658	1.3558	1.356×10^{-3}				
0.2162	1	3.930 × 10 ⁻⁴	0.2520	0.2931	2.931 × 10 ⁻⁴				
550	2544	1	641.19	745.70	0.7457				
0.8578	3.9683	1.560×10^{-3}	1	1.163	1.163×10^{-3}				
0.7376	3.4121	1.341 × 10 ⁻³	0.8598	1	10-3				
737.6	3412.1	1.3410	859.8	1000	1				

ft lbf/lb	Btu/lb	kcal/kg	kg fm/kg	kJ/kg	MJ/kg
1	1.285×10^{-3}	7.139 × 10 ⁻⁴	0.3048	2.989×10^{-3}	2.989 × 10 ⁻⁶
778.2	1	0.5556	237.19	2.326	2.326×10^{-3}
1400.7	1.8	1	426.9	4.187	4.187×10^{-3}
3.281	4.216×10^{-3}	2.342×10^{-3}	1	9.807 × 10 ⁻³	9.807 × 10 ⁻⁶
334.55	0.4299	0.2388	101.97	1	10-3
334553	429.9	238.8	101972	1000	1

Specific Energy (heat/mass) (e.g., calorific value, mass basis, specific latent heat, enthalpy)

Table 10-5 (continued)

Heat Transfer Coefficient (Thermal Conductance) (heat/area × time × degree temperature)

Btu/ft²h°F	Btu/ft2s°F	cal/cm ² s°C	kcal/ft2h°C	W/m²°C	kW/m²°C
1	2.778×10^{-4}	1.356×10^{-4}	0.4536	5.678	5.678×10^{-3}
3600	1	0.4882	1633	20442	20.442
7373	2.048	1	3344	41868	41.868
2.205	6.124 × 10 ⁻⁴	2.990×10^{-4}	1	12.52	0.01252
0.1761	4.892×10^{-5}	2.388×10^{-5}	0.07988	1	10-3
176.1	0.04892	0.02388	79.88	1000	1

	-				
Btu in/ft2h°F	Btu/ft h°F	cal/cm s°C	kcal/m h°C	W/m°C	
1	0.0833	3.445×10^{-4}	0.1240	0.1442	
12	1	4.134×10^{-3}	1.488	1.731	
2903	241.9	1	1 360		
8.064	0.6720	2.778×10^{-3} 1		1.163	
6.933	0.5778	2.388×10^{-3}	2.388×10^{-3} 0.8598		

Thermal Conductivity (heat × length/area × time × degree temperature)

Table 10-5 (continued)

		Stress (force/area)		
dyne/cm ²	N/m ²	lb _f /ft2	kN/m²	lb _t /in ²
1	0.1	2.089×10^{-3}	1 × 10-4	1.45×10^{-5}
10	1	0.02089	1×10^{-3}	1.45×10^{-4}
478.8	47.88	1	0.04788	6.944 × 10 ⁻³
1×10^{4}	1000	20.89	L .	0.145
6.895×10^{4}	6895	144	6.895	1

 $I N/mm^2 = I MN/m^2 = I MPa$

 $1 kg_f/mm^2 = 9.80665 N/mm^2 = 9.80665 MPa$

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Appendix A

Compound	А	В	T _c °C
Fluorine	0.5649	0.2828	-129.0
Chlorine	0.5615	0.2720	144.0
Sulfur dioxide	0.5164	0.2554	157.6
Carbon monoxide	0.2931	0.2706	-140.1
Carbon dioxide	0.4576	0.2590	31.1
Hydrogen chloride	0.4183	0.2619	51.5
Ammonia	0.2312	0.2471	132.4
Water	0.3471	0.2740	374.2
Hydrogen peroxide	0.4375	0.2505	455.0
Hydrogen	0.0315	0.3473	-240.2
Nitrogen	0.3026	0.2763	-146.8
Oxygen	0.4227	0.2797	-118.5
Ethylene	0.2118	0.2784	9.9
Methane	0.1611	0.2877	-82.6
Ethane	0.2202	0.3041	32.3
Propane	0.2204	0.2753	96.7
Benzene	0.3051	0.2714	288.94

Table A1-1 Selected Constants for Liquid Density Equation

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Table A1-1 (continued)							
Toluene	0.2883	0.2624	318.8				
Aniline	0.3392	0.2761	426.0				
Phenol	0.4094	0.3246	420.0				
Cyclopropane	0.2614	0.2826	124.9				
Cyclohexane	0.2729	0.2727	280.3				
1,3 Butadiene	0.2444	0.2710	152.0				
Methanol	0.2928	0.2760	239.4				
Chloroform	0.5165	0.2666	263.4				
Carbon tetrachloride	0.5591	0.2736	283.2				

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Compound	А	В	C × 10-2	D × 10–6	Range °C
Fluorine	-1.576	85.63	-0.04073	-2.725	-219 to 188.1
Chlorine	-0.7681	151.4	-0.08065	0.4075	-101 to 144.0
Sulfur dioxide	-2.67	406.7	0.6141	-12.54	-72.7 to 157.6
Carbon monoxide	-2.346	105.2	0.4613	-19.64	-205 to 140.1
Carbon dioxide	-1.345	21.22	1.034	-34.05	-56.5 to 31.1
Hydrogen chloride	-1.515	194.6	0.3067	-13.76	-114.2 to 51.5
Ammonia	-8.591	876.4	2.681	-36.12	-77.74 to 132.4
Water	-10.73	1828	1.966	-14.66	0.0 to 374.2
Hydrogen peroxide	-1.615	503.8	0.03501	-1.168	-0.43 to 455.0
Hydrogen	-4.857	25.13	14.09	-2773	-259.4 to -240.2
Nitrogen	-12.14	376.1	12.00	-470.9	-209.9 to -195.8
Oxygen	-2.072	93.22	0.6031	-27.21	-218.4 to -118.5

Table A1–2 Selected Constants for Liquid Viscosity Equation

Ethylene	-7.706	468.1	3.725	-76.33	-169.2 to 9.9
Methane	-11.67	499.3	8.125	-226.3	-182.6 to -82.6
Ethane	-4.444	290.1	1.905	-41.64	-183.2 to 32.3
Propane	-3.372	313.5	1.034	-20.26	-187.7 to 96.7
Benzene	2.003	64.66	-1.105	9.648	5.53 to 288.94
Phenol	-8.039	1889	1.055	-6.718	40.75 to 420.0
Cyclopropane	-1.335	116.2	0.01108	-0.03836	93.88 to 238.5
Cyclohexane	-1.910	599.2	-0.06749	0.5026	6.55 to 280.3
1,3 Butadiene	-2.637	434.5	0.1937	-2.907	-108.9 to 152.0
Ethanol	-2.697	700.9	0.2682	-4.917	-105 to 243.1
Chloroform	-1.812	397.5	0.1174	-1.784	63.2 to 263.4
Carbon tetrachloride	-5.658	994.5	1.016	-8.733	-20 to 283.2

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Compound	A	B × 10-3	C × 10-6	D × 10-9	Range °C
Fluorine	-0.288	252.8	-331.0	1464	-219 to -140.0
Chlorine	-0.1322	4.720	-20.37	28.94	-101 to 80.0
Sulfur dioxide	-0.5737	10.34	-40.28	52.85	-72.7 to 150
Carbon monoxide	0.5645	4.798	-143.7	911.95	-205 to -150
Carbon dioxide	-19.30	254.6	-1095.5	1573.3	-56.5 to 20
Hydrogen chloride	-0.1121	7.048	-35.31	66.21	-114.2 to 20
Ammonia	-1.923	31.1	-110.9	137.6	-77.4 to 100
Water	0.6741	2.825	-8.371	8.601	0 to 350
Hydrogen peroxide	0.444	1.199	-2.738	2.615	-0.43 to 425
Hydrogen	3.79	-329.8	12170.9	-2434.8	-259.4 to -245
Nitrogen	-1.064	59.47	-768.7	3357.3	-209.9 to -160
Oxygen	-0.4587	32.34	-395.1	1575.7	-218.4 to -130

Table A1–3 Selected Constants for Liquid Heat Capacitity Equation.

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Ethylene	-0.3402	6.218	-50.12	126.3	-169 to -40
Methane	1.23	-10.33	72.0	-107.3	-182.6 to 110
Ethane	0.1388	8.481	-56.54	126.1	-183.2 to 20
Propane	0.3326	2.332	-13.36	30.16	-187.7 to 80
Benzene	-1.481	15.46	-43.70	44.09	5.53 to 250
Toluene	-0.1461	4.584	-13.46	14.25	-95 to 310
Aniline	0.1407	2.467	-6.085	5.927	-6.15 to 360
Phenol	0.6896	8.218	-18.42	14.47	40.75 to 400
Cyclopropane	-0.02618	6.913	34.77	59.9	-127.42 to 100
Cyclohexane	-1.284	13.39	-35.1	32.27	6.55 to 260
1,3 Butadiene	0.3785	1.049	-5.761	13.74	-108.9 to 120
Methanol	0.8382	-3.231	8.296	-0.1689	-97.6 to 220
Chloroform	-0.09154	3.149	-10.64	12.4	-63.2 to 250
Carbon tetrachloride	-0.01228	2.058	-7.04	8.610	-22.9 to 260

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Compound	А	$B \times 10^{-2}$	C × 10-4	Range °C
Fluorine	612.54	-162.29	-118.41	-219 to -140
Chlorine	559.01	-48.30	-15.24	-101 to 132
Sulfur dioxide	2140.81	-783.66	71.43	-50 to 150
Carbon monoxide	475.48	3.31	-214.26	-205 to -145
Carbon dioxide	972.06	-201.53	-22.99	-56 to 26
Hydrogen chloride	1071.70	-18.44	-65.81	-114 to 31
Ammonia	2551.30	-376.62	-29.35	-77 to 100
Water	-916.62	1254.73	-152.12	0 to 350
Hydrogen peroxide	-466.56	805.86	-87.58	0 to 400
Hydrogen	-20.14	2473.70	-5347.26	-259 to -241
Nitrogen	627.99	-368.91	-22.57	-209 to -152
Oxygen	583.79	-210.49	-48.31	-218 to -135
Ethylene	851.45	-228.94	-4.71	-169.2 to -4.0
Methane	722.72	-144.42	-76.36	182.6 to -90
Ethane	699.31	-165.88	-4.87	-183.2 to 20
Propane	623.51	-126.79	-2.12	-187.7 to 80.0
Benzene	424.26	1.14	-9.03	5.53 to 260
Toluene	485.10	-53.84	-0.59	-95 to 308
Aniline	537.55	-30.42	-1.49	-6.15 to 408
Phenol	440.93	86.70	-7.01	40.75 to 396
Cyclopropane	396.82	-42.10	-6.72	-127.42 to 117
Cyclohexane	388.26	-22.72	-3.30	6.55 to 254
1,3 Butadiene	718.26	-187.18	11.74	-108.9 to 120
Methanol	770.13	-114.28	2.79	-97.6 to 210
Chloroform	390.25	-20.58	-5.06	-63.2 to 237
Carbon tetrachloride	383.95	-45.45	-0.24	-22.9 to 224

 Table A1–4

 Selected Constants for Liquid Thermal Conductivity Equation

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Compound	T₁, °C	T _c , °C	R	Range, °C
Fluorine	-200.0	-129.0	0.8811	-219.6 to -129.0
Chlorine	20.0	144.0	1.0508	-101.0 to 144.0
Sulfur dioxide	30.0	157.6	1.1768	-72.7 to 157.6
Carbon monoxide	-193.0	-140.1	1.1441	-191.5 to -140.1
Carbon dioxide	20.0	31.1	1.3015	-56.5 to 31.1
Hydrogen chloride	-93.0	51.5	1.0972	-114.2 to 51.5
Ammonia	-45.0	132.4	1.1548	-77.74 to 132.4
Water	25.0	374.2	0.8105	0 to 100.0
Water	100.0	374.2	1.1690	100.0 to 374.2
Hydrogen peroxide	18.2	455.0	0.9141	-0.43 to 455.0
Hydrogen	-256.0	-240.2	1.1012	-259 to -240.2
Nitrogen	-203.0	-146.8	1.2123	-209.9 to -146.8
Oxygen	-202.0	-118.5	1.1933	-218.4 to -118.5
Ethylene	-120.0	9.9	1.2760	-169.2 to 9.9
Methane	-168.16	-82.6	1.3941	-182.6 to -82.6
Ethane	-120.0	32.3	1.2060	-183.2 to 32.3
Propane	-90.0	96.7	1.1982	-187.7 to 96.7
Benzene	20.0	288.94	1.2243	5.53 to 288.94
Touluene	20.0	318.8	1.2364	-95.0 to 318.8
Aniline	20.0	426.0	1.1022	-6.15 to 426.0
Phenol	60.0	420.0	1.0725	40.75 to 420.0
Cyclopropane	10.0	124.9	1.3201	-127.42 to 124.9
Cyclohexane	20.0	280.3	1.4246	6.55 to 280.3
1,3 Butadiene	40.0	152.0	1.2055	-4.41 to 152.0
Methanol	20.0	239.4	0.8115	-97.6 to 239.4
Chloroform	25.0	263.4	1.1824	-63.2 to 263.4
Carbon tetrachloride	30.0	283.2	1.2278	-22.9 to 283.2

 Table A1–5

 Selected Constants for Liquid Surface Tension Equation

By Permission: C.L. Yaws et al., Physical Properties, A Chemical Engineering publication, McGraw-Hill, New York, 1977.

Compound	А	В	С	D × 10-3	E × 10-6	Range °C
Fluorine	21.480	-516.51	-7.1218	14.355		-219.2 to -129.0
Chlorine	42.262	-2009.8	-13.963	9.3705	_	-100.4 to 144.1
Sulfur dioxide	46.554	-2456.3	-15.169	9.0026	_	-67.6 to 157.6
Carbon monoxide	32.863	-606.91	-12.969	27.551	_	-200.7 to -140
Carbon dioxide	47.544	-1792.2	-16.559	13.833	_	-56.0 to 31.2
Hydrogen chloride	136.05	-3047.3	-58.416	95.496	-58.507	150.8 to 51.5
Ammonia	38.44	-2066.2	-12.105	7.7768		-65 to 132.4
Water	16.373	-2818.6	-1.6908	-5.7546	4.0073	0.0 to 374.2
Hydrogen peroxide	44.791	-4022.7	-13.076	4.5627	_	-0.01 to 455
Hydrogen	5.2366	-46.28	-0.44809	25.290		-259.4 to -240.2
Nitrogen	21.623	-455.57	-7.5107	17.214		-210 to -146.8
Oxygen	5.6486	-411.30	1.8118	-25.042	62.612	-219.1 to -118.4

Table A1–6 Selected Constants for Liquid Vapor Pressure Equation

814

Ethylene	30.895	-1196.8	-10.153	9.9351		-169 to 10.1
Methane	22.573	-656.24	-7.3942	11.896		-182.2 to 81.9
Ethane	16.316	-1074.8	-3.1434	4.5534	10.373	-179.5 to 32.3
Propane	36.007	-1737.2	-11.666	8.5187		-128.9 to 96.8
Benzene	51.204	-3245.7	-16.403	7.540		7.6 to 289.4
Toluene	115.21	-4918.1	-43.467	38.548	-13.496	-60.0 to 320.6
Aniline	18.893	-3104.0	-3.4714	0.02743		77.4 to 426
Phenol	672.71	-22197.3	266.67	226.43	-73.731	67.8 to 420
Cyclopropane	38.45	-1865.2	-12.578	8.9375		-90 to 125.2
Cyclohexane	64.753	-3619.2	-21.753	10.742		-25.4 to 280.3
1,3 Butadiene	41.401	-2181.2	-13.451	8.4524	—	-108.9 to 152
Methanol	-42.629	-1186.2	23.279	-35.082	17.578	-67.4 to 240
Chloroform	26.828	-2292.6	-7.186	3.1365		-42 to 263
Carbon tetrachloride	50.612	-3135.7	-16.313	7.8036		-69.7 to 283.2

By Permission: C.L. Yaws et al., Physical Properties, A Chemical Engineering publication, McGraw-Hill, New York, 1977.

Compound	∆H _{V1} , cal/g	T₁, °C	T _c , °C	R
Fluorine	41.1	-188.1	-129.0	0.38
Chlorine	69.6	-34.06	144.0	0.38
Sulfur dioxide	93.0	-10.0	157.6	0.38
Carbon monoxide	51.6	-191.5	-140.1	0.38
Carbon dioxide	56.1	0.0	31.1	0.38
Hydrogen chloride	105.8	-85.03	51.5	0.38
Ammonia	327.4	-33.43	132.4	0.38
Water	538.7	100.0	374.2	0.38
Hydrogen peroxide	321.9	150.2	455.0	0.38
Hydrogen	107.0	-252.8	-240.2	0.237
Nitrogen	47.5	-195.8	-146.8	0.38
Oxygen	50.9	-183.0	-118.5	0.38
Ethylene	115.4	-103.7	9.9	0.38
Methane	121.7	-161.5	-82.6	0.38
Ethane	116.7	-88.2	32.3	0.38
Propane	101.8	-42.1	96.7	0.38
Benzene	94.1	80.1	288.94	0.38
Toluene	86.1	110.6	318.8	0.38
Aniline	112.4	184.4	426.0	0.38
Phenol	116.4	181.8	420.0	0.38
Cyclopropane	113.8	-32.8	124.9	0.38
1,3 Butadiene	100.2	-4.41	152.0	0.38
Methanol	260.1	64.7	239.4	0.40
Chloroform	58.9	61.3	263.4	0.38
Carbon tetrachloride	46.55	76.7	283.2	0.38

 Table A1–7

 Selected Constants for Enthalpy of Vaporization Equation

By Permission: C.L. Yaws et al., Physical Properties, A Chemical Engineering publication, McGraw-Hill, New York, 1977.

Compound	Α	В×10−3	C × 10-6	D × 10-9	Range K
Fluorine	5.89	6.94	-5.48	1.52	298–1500
Chlorine	7.00	5.05	-4.39	1.30	298-1500
Sulfur dioxide	5.85	15.40	-11.1	2.91	298-1500
Carbon monoxide	6.92	-0.65	2.80	-1.14	298-1500
Carbon dioxide	5.14	15.4	-9.94	2.42	298-1500
Hydrogen chloride	7.24	-1.76	3.07	-1.00	298-1500
Ammonia	6.07	8.23	-0.16	-0.66	298-1500
Water	8.10	-0.72	3.63	-1.16	298-1500
Hydrogen peroxide	5.52	19.80	-13.90	3.74	298-1500
Hydrogen	6.88	-0.022	0.21	0.13	298-1500
Nitrogen	7.07	-1.32	3.31	-1.26	298-1500
Oxygen	6.22	2.71	-0.37	-0.22	298–1500
Ethylene	0.934	36.9	-19.3	4.01	298-1500
Methane	5.04	9.32	8.87	-5.37	298-1500
Ethane	2.46	36.1	-7.0	-0.46	298-1500
Propane	-0.58	69.9	-32.9	6.54	298-1500
Benzene	-8.79	116.0	-76.1	18.9	298-1500
Toluene	-9.34	138.5	-87.2	20.6	298-1500
Aniline	-8.11	143.4	-107.0	30.7	298-1500
Phenol	-5.68	126.0	-85.4	20.5	298-1500
Cyclopropane	-6.28	79.8	-50.5	12.2	298-1500
1,3 Butadiene	-0.56	81.1	-53.5	13.6	298-1500
Methanol	3.62	24.9	-7.05		298–1500
Chloroform	7.71	34.3	-26.40	7.29	298–1500
Carbon tetrachloride	12.6	33.2	-29.3	8.58	298-1500

 Table A1–8

 Selected Constants for Gas Heat Capacity Equation

By Permission: C. L. Yaws et al. Physical Properties, A Chemical Engineering publication, McGraw-Hill, New York, 1977.

Compound	А	B × 10-3	C × 10-6	Range, K
Sulfur dioxide	-69.6	-5.29		298 to 717
Sulfur dioxide	-86.9	0.32		717 to 1500
Nitrogen dioxide	7.94	-0.188		298 to 1500
Carbon monoxide	-26.5	0.84	-1.05	298 to 1500
Carbon dioxide	-93.9	-0.43	—	298 to 1500
Hydrogen chloride	-21.9	-0.610	—	298 to 1500
Ammonia	-9.34	-6.20	2.36	298 to 1500
Water	-57.4	-1.79	—	298 to 1500
Hydrogen peroxide	-31.8	-3.04	1.19	298 to 1500
Methane	-15.4	-9.59	3.50	298 to 1500
Ethane	-16.4	-14.8	6.13	298 to 1500
Propane	-20.0	-19.1	8.15	298 to 1500
Benzene	23.7	-15.3	6.27	298 to 1500
Toluene	16.8	-19.0	7.84	298 to 1500
Aniline	25.2	-17.6	8.98	298 to 1500
Phenol	-19.3	-14.6	7.18	298 to 1500
Cyclopropane	16.9	-16.7	7.90	298 to 1500
Cyclohexane	-21.6	-32.0	15.80	298 to 1500
1,3 Butadiene	29.0	-10.1	4.13	298 to 1500
Methanol	-44.96	-11.9	4.98	298 to 1500
Chloroform	-24.7	0.0334		298 to 1500
Carbon tetrachloride	-24.0	2.42		298 to 1500

 Table A1–9

 Selected Constants for Heat of Formation Equation

By Permission: C. L. Yaws et al. Physical Properties, A Chemical Engineering publication, McGraw-Hill, New York, 1977.

Compound	А	$B \times 10^{-3}$	Range, K
Sulfur dioxide	-71.9	0.25	298 to 717
Sulfur dioxide	-86.8	17.7	717 to 1500
Nitrogen dioxide	7.82	15.1	298 to 1500
Carbon monoxide	-26.5	-21.3	298 to 1500
Carbon dioxide	-94.2	-0.42	298 to 1500
Hydrogen chloride	-22.3	-1.72	298 to 1500
Ammonia	-12.3	27.2	298 to 1500
Water	-58.6	12.7	298 to 1500
Hydrogen peroxide	-33.2	26.4	298 to 1500
Methane	-20.1	24.9	298 to 1500
Ethane	-23.3	49.7	298 to 1500
Propane	-28.8	74.7	298 to 1500
Benzene	16.7	45.9	298 to 1500
Toluene	7.8	68.7	298 to 1500
Aniline	18.5	70.6	298 to 1500
Phenol	-25.0	56.5	298 to 1500
Cyclopropane	10.3	47.7	298 to 1500
Cyclohexane	-35.1	139.0	298 to 1500
1,3 Butadiene	24.2	38.6	298 to 1500
Methanol	-50.2	36.5	298 to 1500
Chloroform	-24.8	26.9	298 to 1500
Carbon tetrachloride	-22.6	32.1	298 to 1500

 Table A1–10

 Selected Constants for Free Energy of Formation Equation

By Permission: C. L. Yaws et al. Physical Properties, A Chemical Engineering publication, McGraw-Hill, New York, 1977.

Compound	А	B × 10-2	C × 10-4	D × 10-8	Range °C
Fluorine	1.8654	19.79	1.24	-17.77	-130 to 525
Chlorine	3.25	5.8	0.21	-1.25	-80 to 1200
Sulfur dioxide	-19.31	15.15	-0.33	0.55	0 to 1400
Carbon monoxide	1.21	21.79	-0.8416	1.958	-160 to 1400
Carbon dioxide	-17.23	19.14	0.1308	-2.514	-90 to 1400
Hydrogen chloride	-0.26	12.67	-0.25	0.16	-150 to 1400
Ammonia	0.91	12.87	2.93	-8.68	0 to 1400
Water	17.53	-2.42	4.3	-21.73	0 to 800
Hydrogen peroxide	-21.07	16.97	0.17	-1.56	0 to 1200
Hydrogen	19.34	159.74	-9.93	37.29	-160 to 1200
Nitrogen	0.9359	23.44	-1.21	3.591	-160 to 1200
Oxygen	-0.7816	23.8	0.8939	2.324	-160 to 1200
Ethylene	-42.04	28.65	0.7963	-3.262	-75 to 1000
Methane	-4.463	20.84	2.815	-8.631	0 to 1000
Ethane	-75.8	52.57	-4.593	39.74	0 to 750
Propane	4.438	-1.122	5.198	-20.08	0 to 1000
Benzene	-20.19	8.64	2.34	-9.69	0 to 1000
Toluene	18.14	-9.57	5.66	-22.22	0 to 1000
Aniline	-26.39	11.89	1.55	-4.30	0 to 1000
Phenol	-31.87	15.26	1.74	-4.40	0 to 1000
Cyclopropane	-20.46	9.74	3.77	-16.28	0 to 800
Cyclohexane	-20.57	4.45	4.07	-17.31	0 to 800
1,3 Butadiene	-67.92	29.96	1.74	-12.20	0 to 1000
Methanol	-18.62	9.95	2.90	-12.38	0 to 1000
Chloroform	-5.732	6.29	0.5904	-3.352	25 to 1000
Carbon tetrachloride	-0.4161	4.067	0.6115	-3.566	25 to 1000

 Table A1–11

 Selected Constants for Gas Thermal Conductivity Equation

By Permission: C. L. Yaws et al. Physical Properties, A Chemical Engineering publication, McGraw-Hill, New York, 1977.

Compound	Α	B × 10-2	C × 10-6	Range °C
Fluorine	22.09	76.9	-211.6	-200 to 1000
Chlorine	5.175	45.69	-88.54	-200 to 1200
Sulfur dioxide	-3.793	46.45	-72.76	-100 to 1400
Carbon monoxide	32.28	47.47	-96.48	-200 to 1400
Carbon dioxide	25.45	45.49	-86.49	-100 to 1400
Hydrogen chloride	-9.554	54.45	-96.56	-120 to 1400
Ammonia	-9.372	38.99	-44.05	-200 to 1200
Water	-31.89	41.45	-8.272	0 to 1000
Hydrogen peroxide	5.381	28.98	38.40	-80 to 1000
Hydrogen	21.87	22.2	-37.51	-160 to 1200
Nitrogen	30.43	49.89	-109.3	-160 to 1200
Oxygen	18.11	66.32	-187.9	-160 to 1000
Ethylene	3.586	35.13	-80.55	-100 to 800
Methane	15.96	34.39	-81.40	-80 to 1000
Ethane	5.576	30.64	-53.07	-80 to 1000
Propane	4.912	27.12	-38.06	-80 to 1000
Benzene	-15.76	32.45	-72.32	0 to 1000
Toluene	-8.421	27.11	-40.18	0 to 1000
Aniline	-14.98	29.03	-1.116	0 to 1000
Phenol	-16.41	32.00	0	0 to 1000
Cyclopropane	-7.787	34.78	-81.3	0 to 1000
Cyclohexane	-4.705	26.32	-44.1	0 to 1000
1,3 Butadiene	-10.67	34.32	-80.8	0 to 1000
Methanol	-5.636	34.45	-3.34	0 to 900
Chloroform	6.688	37.26	-50.87	0 to 1000
Carbon tetrachloride	5.698	32.73	-40.28	0 to 1000

Table A1–12Selected Constants for Gas Viscosity Equation

By Permission: C. L. Yaws et al. Physical Properties, A Chemical Engineering publication, McGraw-Hill, New York, 1977.

Appendix B

Table B-1 Compressibility Factor of Natural Gas

****	**********
TEMPERATURE, oF	SPECIFIC GRAVITY
60.	,550
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.98160
200.	.97934
300.	.97263
400.	.96270
500.	.95042
600.	.93651
700.	.92156
800.	.90610
900.	.89053
1000.	.87520
1100.	.86041
1200.	.84636
1300.	.83322
1400.	.82110
1500.	.81008
1600.	.80021
1700.	.79151
1800.	.78399
1900.	.77767
2000.	.77253
2100.	.76860
2200.	.76586
2300.	.76432
2400.	.76395
2500.	.76473

Table B-1 (continued)

*******		*************	***********************
	RAVITY	SPECIFIC G	TEMPERATURE, oF
		.550	60.
Z	FACTOR	COMPRESSIBILITY	PRESSURE psia.
		.76663	2600.
		.76960	2700.
		.77354	2800.
		.77838	2900.
		.78401	3000.
		.79032	3100.
		.79718	3200.
		.80447	3300.
		.81208	3400.
		.81989	3500.
		.82780	3600.
		.83573	3700.
		.84363	3800.
		.85144	3900.
		.85916	4000.
		.86678	4100.
		.87436	4200.
		.88195	4300.
		.88966	4400.
		.89761	4500.
		.90597	4600.
		.91495	4700.
		.92478	4800.
		.93575	4900.
		.94817	5000.

Table B-2			
Compressibility	Factor o	of Natural	Gas

TEMPERATURE, OF 70.	SPECIFIC GRAVITY .550
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.98208
200.	.98046
300.	.97453
400.	.96546
500.	.95411
600.	.94117
700.	.92721
800.	.91272
900.	.89810
1000.	.88369
1100.	.86975
1200.	.85649
1300.	.84406
1400.	.83257
1500.	.82209
1600.	.81267
1700.	.80432
1800.	.79706
1900.	.79089
2000.	.78583
2100.	.78187
2200.	.77903
2300.	.77729
2400.	.77665
2500.	.77710
2600.	.77859
2700.	.78109
2800.	.78452
2900.	.78880
3000.	.79383
3100.	.79951
3200.	.80572
3300.	.81234
3400.	.81926
3500.	.82638
3600.	.83360
3700.	.84085
3800.	.84806
3900.	.85519
4000.	.86223
4100.	.86919
4200.	.87610
4300.	.88303
4400.	.89006
4500.	.89733
4600.	.90499
4700.	.91323
4800.	.92228
4900.	.93241
5000.	.94392

Table B-3				
Compressibility	Factor	of	Natural	Gas

TEMPERATURE, oF 80.	SPECIFIC GRAVITY .550
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.98250
200.	.98147
300.	.97625
400.	.96798
500.	.95750
600.	.94547
700.	.93243
800.	.91886
900.	.90513
1000.	.89158
1100.	.87845
1200.	.86594
1300.	.85419
1400.	.84330
1500.	.83334
1600.	.82434
1700.	.81634
1800.	.80933
1900.	.80333
2000.	.79834
2100.	.79437
2200.	.79143
2300.	.78952
2400.	.78864
2500.	.78878
2600.	.78990
2700.	.79197
2800.	.79492
2900.	.79868
3000.	.80317
3100.	.80826
3200.	.81387
3300.	.81987
3400.	.82616
3500.	.83264
3600.	.83922
3700.	.84582
3800.	.85239
3900.	.85889
4000.	.86531
4100.	.87165
4200.	.87794
4300.	.88425
4400.	.89066
4500.	.89729
4600.	.90429
4700.	.91185
4800.	.92017
4900.	.92952
5000.	.94017

**************************************	**************************************
90. PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.98288
200.	.98238
300.	.97781
400.	.97029
500.	.96061
600.	.94943
700.	.93726
800.	.92455
900.	.91167
1000.	.89893
1100.	.88657
1200.	.87476
1300.	.86366
1400.	.85334
1500.	.84387
1600.	.83529
1700.	.82762
1800.	.82085
1900.	.81501
2000.	.81010
2100.	.80613
2200.	.80311
2300.	.80105
2400.	.79995
2500.	.79980
2600.	.80058
2700.	.80226
2800.	.80477
2900.	.80806
3000.	.81203
3100.	.81658
3200.	.82163
3300.	.82705
3400.	.83275
3500.	.83864
3600.	.84462
3700.	.85063
3800.	.85661
3900.	.86253
4000.	.86836
4100.	.87413
4200.	.87984
4300.	.88558
4400.	.89141
4500.	.89745
4600.	.90385
4700.	.91076
4800.	.91841
4900.	.92702
5000.	.93688

Table B-4Compressibility Factor of Natural Gas

Ta	able B-5		
Compressibility	Factor of	Natural	Gas

**************************************	**************************************
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.98321
200.	.98320
300.	.97923
400.	.97239
500.	.96348
600.	.95308
700.	.94173
800.	.92983
900.	.91775
1000.	.90577
1100.	.89413
1200.	.88300
1300.	.87251
1400.	.86274
1500.	.85374
1600.	.84556
1700.	.83820
1800.	.83167
1900.	.82599
2000.	.82116
2100.	.81720
2200.	.81411
2300.	.81192
2400.	.81062
2500.	.81021
2600.	.81067
2700.	.81199
2800.	.81410
2900.	.81694
3000.	.82043
3100.	.82449
3200.	.82901
3300.	.83390
3400.	.83905
3500.	.84439
3600.	.84982
3700.	.85528
3800.	.86071
3900.	.86608
4000.	.87137
4100.	.87660
4200.	.88179
4300.	.88699
4400.	.89228
4500.	.89778
4600.	.90360
4700.	.90993
4800.	.91694
4900.	.92487
5000.	.93398

Table B-6Compressibility Factor of Natural Gas

TEMPERATURE, oF 60.	SPECIFIC GRAVITY .600
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.96443
200.	.95901
300.	.94996
400.	.93810
500.	.92410
600.	.90857
700.	.89204
800.	.87496
900.	.85775
1000.	.84077
1100.	.82432
1200.	.80866
1300.	.79403
1400.	.78060
1500.	.76851
1600.	.75787
1700.	.74877
1800.	.74124
1900.	.73532
2000.	.73098
2100.	.72821
2200.	.72695
2300.	.72715
2400.	.72871
2500.	.73155
2600. 2700.	.73556 .74062
2800.	.74663
2900.	.75345
3000.	.76096
3100.	.76903
3200.	.77756
3300.	.78642
3400.	.79552
3500.	.80475
3600.	.81403
3700.	.82330
3800.	.83251
3900.	.84163
4000.	.85066
4100.	.85961
4200.	.86852
4300.	.87748
4400.	.88658
4500.	.89598
4600.	.90584
4700.	.91638
4800.	.92786
4900.	.94058
5000.	.95488

Table B-7			
Compressibility Factor of Natural Gas			

*****	*******
TEMPERATURE, OF	SPECIFIC GRAVITY
70.	.600
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.96511
200.	.96051
300.	.95238
400.	.94152
500.	.92858
600.	.91413
700.	.89868
800.	.88267
900.	.86649
1000.	.85049
1100.	.83495
1200.	.82015
1300.	.80628
1400.	.79352
1400.	.78201
1600.	.77186
1700.	.76314
1800.	.75589
1900.	.75015
2000.	.74589
2100.	.74311
2200.	.74175
2300.	.74175
2400.	.74304
2500.	.74553
2600.	.74912
2700.	.75370
2800.	.75917
2900.	.76540
3000.	.77229
3100.	.77970
3200.	.78754
3300.	.79569
3400.	.80406
3500.	.81255
3600.	.82110
3700.	.82963
3800.	.83810
3900.	.84649
4000.	.85478
4100.	.86300
4200.	.87119
4300.	.87942
4400.	.88779
4500.	.89643
4600.	.90552
4700.	.91525
4800.	.92587
4900.	.93767
5000.	.95097
*	

TEMPERATURE, oF 80.	SPECIFIC GRAVITY .600
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.96572
200.	.96186
300.	.95459
400.	.94466
500.	.93271
600.	.91927
700.	.90483
800.	.88983
900.	.87462
1000.	.85955
1100.	.84490
1200.	.83090
1300.	.81776
1400.	.80565
1500.	.79470
	.78502
1600.	
1700.	.77667
1800.	.76971
1900.	.76415
2000.	.75999
2100.	.75721
2200.	.75576
2300.	.75560
2400.	.75665
2500.	.75882
2600.	.76203
2700.	.76618
2800.	.77115
2900.	.77684
3000.	.78315
3100.	.78995
3200.	.79716
3300.	.80465
3400.	.81235
3500.	.82016
3600.	.82802
3700.	.83587
3800.	.84366
3900.	.85136
4000.	.85898
4100.	.86653
4200.	.87404
4300.	.88160
4400.	.88928
4500.	.89723
4600.	.90560
4700.	.91458
4800.	.92440
4900.	.93534
	• 7 3 3 3 4

Table B-8Compressibility Factor of Natural Gas

TEMPERATURE, oF 90.	SPECIFIC GRAVITY .600
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.96626
200.	.96308
300.	.95660
400.	.94754
500.	.93650
600.	.92401
700.	.91054
800.	.89648
900.	.88220
1000.	.86801
1100.	.85419
1200.	.84096
1300.	.82852
1400.	.81703
1500.	.80662
1600.	.79739
1700.	.78941
1800.	.78273
1900.	.77735
2000.	.77330
2100.	.77053
2200.	.76902
2300.	.76872
2400.	.76955
2500.	.77144
2600.	.77431
2700.	.77805
2800.	.78257
2900.	.78777
3000.	.79354
3100.	.79979
3200.	.80640
3300.	.81329
3400.	.82037
3500.	.82756
3600.	.83478
3700.	.84199
3800.	.84915
3900.	.85622
4000.	.86322
4100.	.87014
4200.	.87703
4300.	.88396
4400.	.89102
4500.	.89832
4600.	.90602
4700.	.91429
4800.	.92337
4900.	.93351
5000.	.94499

Table B-9 Compressibility Factor of Natural Gas

Table B-10			
Compressibility Factor of Natural Gas			

**************************************	SPECIFIC GRAVITY
100.	.600
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.96674
200.	.96419
300.	.95843
400.	.95018
500.	.94000
600.	.92840
700.	.91583
800.	.90266
900.	.88926
1000.	.87590
1100.	.86287
1200.	.85037
1300.	.83860
1400.	.82771
1500.	.81782
1600.	.80903
1700.	.80141
1800.	.79499
1900.	.78981
2000.	.78586
2100.	.78312
2200.	.78156
2300.	.78113
2400.	.78178
2500.	.78341
2600.	.78596
2700.	.78934
2800.	.79345
2900.	.79819
3000.	.80348
3100.	.80920
3200.	.81528
3300.	.82161
3400.	.82811
3500.	.83472
3600.	.84136
3700.	.84798
3800.	.85455
3900.	.86104
4000.	.86745
4100.	.87380
4200.	.88012
4300.	.88647
4400.	.89293
4500.	.89964
4600.	.90672
4700.	.91434
4800.	.92273
4900.	.93212
5000.	.94279
5000.	

Table B-11				
Compressibility	Factor	of	Natural	Gas

*****	***************************************
TEMPERATURE, oF	SPECIFIC GRAVITY
60.	.650
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.94835
200.	.93926
300.	.92722
400.	.91276
500.	.89642
600.	.87870
700.	.86004
800.	.84088
900.	.82161
1000.	.80258
1100.	.78414
1200.	.76659
1300.	.75021 .73524
1400. 1500.	.73524
1600.	.71033
1700.	.70067
1800.	.69302
1900.	.68738
2000.	.68375
2100.	.68206
2200.	.68223
2300.	.68411
2400.	.68756
2500.	.69242
2600.	.69851
2700.	.70565
2800.	.71370
2900.	.72248
3000.	.73185
3100.	.74169
3200.	.75188
3300.	.76231
3400.	.77288 .78353
3500. 3600.	.78353
3700.	.80478
3800.	.81531
3900.	.82573
4000.	.83606
4100.	.84632
4200.	.85656
4300.	.86688
4400.	.87737
4500.	.88819
4600.	.89954
4700.	.91163
4800.	.92474
4900.	.93919
5000.	.95536

70. .650 PRESSURE psia. COMPRESSIBILITY FACTOR Z 100. .94926 200. .94120 300. .93025 400. .91636 500. .90180 600. .86527 700. .86780 800. .64793 900. .61365 1100. .79618 1200. .74959 1400. .73694 1600. .73694 1600. .70926 1900. .700381 2000. .70027 2100. .69856 2200. .70794 2500. .70794 2500. .72769 3000. .72358 3200. .72354 3000. .72354 3000. .72354 3000. .72354 3000. .72354 3000. .72354 3000. .72354 3000. .7235	TEMPERATURE, oF	SPECIFIC GRAVITY
100. .94926 200. .94120 300. .91025 400. .91696 500. .90180 600. .88527 700. .86780 800. .84979 900. .81365 1100. .79618 1200. .77953 1300. .76395 1400. .73644 1600. .73588 1700. .71663 1800. .70926 1900. .70381 2000. .70027 2100. .69856 2200. .69856 2200. .7029 2400. .73584 3000. .72052 2800. .72759 3300. .72052 2800. .72584 3000. .72622 2800. .72759 3300. .72623 3400. .72829 3500. .72584	70.	.650
100. .94926 200. .94120 300. .93025 400. .91696 500. .90180 600. .85780 800. .84979 900. .83162 1000. .81365 1100. .77953 1300. .76395 1400. .73694 1600. .72588 1700. .71663 1800. .70926 1900. .70381 2000. .69856 2200. .69856 2200. .69856 2200. .69856 2200. .7045 2500. .70794 2600. .72769 2900. .73584 3000. .74455 3100. .75368 3200. .76313 3300. .727280 3400. .79245 3600. .82183 3900. .82145 3000. .7259 3500. .79245 <th>PRESSURE psia.</th> <th></th>	PRESSURE psia.	
300. 93025 400. 91696 500. 90180 600. .86780 800. .84979 900. .83162 1000. .79618 1200. .77693 1400. .73694 1600. .72588 1700. .70644 1600. .70926 1900. .70381 2000. .70027 2100. .69956 2200. .69956 2200. .69956 2200. .7024 2600. .70794 2600. .72769 2900. .73584 3000. .74513 3100. .72568 3200. .72769 2900. .73584 3000. .7455 3100. .72568 3200. .72769 3900. .72584 3000. .7455 3100. .72586 3200. .72580 3600. .80230	100.	
400. 91696 500. 90180 600. 88527 700. 86780 800. 84979 900. 81365 1100. 79618 1200. 77953 1300. 77953 1300. 77953 1400. 77953 1500. 773694 1600. 72588 1700. 701663 1800. 70926 1900. 70381 2000. 70027 2100. 69856 2200. 69861 2300. 70145 2500. 70794 2600. 73584 3000. 7455 3100. 77269 2900. 73584 3000. 77280 3400. 78259 3500. 72769 3900. 81210 3800. 82183 3900. 82183 3900. 81210 3800. 82183 3900. </td <td></td> <td>.94120</td>		.94120
500. 90180 600. .88527 700. .86780 800. .84979 900. .81362 1000. .7953 1100. .79618 1200. .71663 1600. .72588 1700. .70694 1600. .72588 1700. .70631 1800. .70381 2000. .669861 2000. .70027 2100. .669861 2200. .70029 2400. .70794 2600. .71358 2700. .72536 3100. .72545 3100. .72545 3100. .72546 3200. .72536 3200. .72536 3200. .72536 3200. .72545 3600. .82183 3900. .82183 3900. .82183 3900. .82183		.93025
600. .88527 700. .86780 800. .84979 900. .81362 1000. .81365 1100. .79618 1200. .76395 1300. .76395 1400. .74969 1500. .73694 1600. .72588 1700. .71663 1800. .70926 1900. .70027 2100. .69856 2200. .69861 2300. .70794 2600. .71358 3000. .72769 2900. .73584 3000. .74455 3100. .75368 3200. .72769 2900. .73584 3000. .74455 3100. .7268 3200. .72845 3600. .80230 3700. .82183 3900. .83145 4000. .85945 4100. .85945 4100. .85945		.91696
700. .86780 800. .83162 1000. .81365 1100. .79618 1200. .77953 1300. .76395 1400. .74969 1500. .72588 1700. .71663 1800. .70926 1900. .70381 200. .69856 2200. .69861 2300. .70029 2400. .70345 2500. .70794 2600. .72588 2700. .7269 2800. .72769 2800. .72769 2900. .73584 3000. .74455 3100. .725368 3200. .72769 2800. .72769 3900. .78259 3500. .79245 3600. .82183 3900. .82183 3900. .83145 4000. .82183 3900. .83145 4000. .85941		
800.		
900. .83162 1000. .81365 1100. .79618 1200. .77953 1300. .76395 1400. .74969 1500. .72588 1770. .71663 1800. .70926 1900. .70381 2000. .70027 2100. .69861 2300. .70029 2400. .70345 2500. .70794 2600. .71358 2700. .72022 2800. .72769 2900. .73584 3000. .77455 3100. .75368 3200. .77513 3300. .77280 3400. .78259 3500. .81210 3800. .8145 4000. .85045 4200. .85045 4200. .85950 4300. .85910 4400. .85910 4500. .88910 4500. .89959		
1000. 81365 1100. .79618 1200. .77953 1300. .76395 1400. .74969 1500. .73694 1600. .72588 1700. .71663 1800. .70926 1900. .70381 2000. .69856 2200. .69861 2300. .70145 2500. .70794 2600. .71358 2700. .72568 200. .73654 2500. .70794 2600. .71358 2700. .72769 2900. .73584 3000. .75368 3200. .76313 3300. .72280 3400. .78259 3500. .79245 3600. .82183 3900. .81145 4000. .8595 4100. .85910 4300. .85959		
1100. .79618 1200. .77953 1300. .76395 1400. .74969 1500. .7364 1600. .72588 1700. .71663 1800. .70926 1900. .70381 2000. .70027 2100. .669861 2300. .70029 2400. .70345 2500. .70794 2600. .71358 2700. .7269 2800. .72769 2900. .73584 3000. .75368 3200. .76313 3300. .77280 3400. .78259 3500. .79245 3600. .82183 3900. .81210 3800. .82183 3900. .81210 3800. .82183 3900. .8145 4000. .85991 4100. .85990 4200. .85990 4300. .89590		
1200. .77953 1300. .76395 1400. .74969 1500. .73694 1600. .72588 1700. .71663 1800. .70926 1900. .70381 2000. .69856 2200. .69861 2300. .70029 2400. .70345 2500. .70794 2600. .72588 2700. .72022 2800. .72769 2900. .73584 3000. .74455 3100. .725368 3200. .725368 3200. .725368 3200. .725368 3200. .725368 3200. .7280 3400. .82193 3900. .82183 3900. .82183 3900. .82183 3900. .83145 4000. .85990 4300. .85990 4300. .85990 4300. .85990		
1300. .76395 1400. .74969 1500. .7354 1600. .72588 1700. .71663 1800. .70926 1900. .70381 2000. .70027 2100. .69866 2200. .69861 2300. .70794 2600. .71358 2700. .72769 2800. .72769 2900. .73584 3000. .74455 3100. .75368 3200. .7280 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .82183 3900. .8145 4000. .86941 4100. .85045 4200. .85990 4300. .88910 4500. .88910 4600. .89959 4700. .81079 4800. .92296 4700. .91079		
1400. .74969 1500. .73694 1600. .72588 1700. .71663 1800. .70926 1900. .70381 2000. .69856 2200. .69861 2300. .70029 2400. .70345 2500. .70794 2600. .71358 2700. .72769 2900. .73584 3000. .74455 3100. .75368 3200. .76313 3300. .77280 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .8145 4000. .85945 4200. .85990 4300. .85990 4300. .85910 4500. .88910 4500. .88910 4500. .88910 4500. .92296 4700. .91079 4800. .92296		
1500. .73694 1600. .72588 1700. .71663 1800. .70926 1900. .70381 2000. .70027 2100. .69861 2300. .70345 2500. .70794 2600. .71358 2700. .7269 2800. .72769 2900. .73584 3000. .74455 3100. .77269 2900. .73584 3000. .74455 3100. .77280 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .81210 3800. .81245 4000. .85045 4200. .85991 4100. .85045 4200. .85991 4300. .85910 4500. .88910 4500. .88910 4500. .93079 4700. .91079		
1600. .72588 1700. .71663 1800. .70926 1900. .70381 2000. .69856 2200. .69861 2300. .70029 2400. .70345 2500. .70794 2600. .71358 2700. .72769 2900. .73584 3000. .74455 3100. .75368 3200. .76313 3300. .77280 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .81210 3800. .81210 3800. .81216 3900. .81217 3900. .8145 4000. .85990 4100. .85941 4400. .87910 4500. .89959 4700. .91079 4800. .92296		
1700. .71663 1800. .70326 1900. .70331 2000. .69856 2200. .69861 2300. .70029 2400. .70345 2500. .70794 2600. .71358 2700. .72022 2800. .72769 2900. .73584 3000. .74455 3100. .75368 3200. .72780 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .81283 3900. .81245 4000. .84098 4100. .85045 4200. .85990 4300. .85941 4400. .87910 4500. .89959 4700. .91079 4800. .92296		
1800. .70926 1900. .70031 2000. .70027 2100. .69856 2200. .69861 2300. .70029 2400. .70345 2500. .70794 2600. .71358 2700. .72022 2800. .72769 2900. .73584 3000. .74455 3100. .75368 3200. .76313 3300. .77280 3400. .78259 3500. .79245 3600. .81210 3800. .81145 4000. .84098 4100. .85045 4200. .85990 4300. .85990 4300. .85910 4600. .89959 4700. .89109 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
1900. .70381 2000. .70027 2100. .69856 2200. .69861 2300. .70029 2400. .70794 2600. .71358 2700. .72022 2800. .72769 2900. .73584 3000. .74455 3100. .77368 3200. .76313 3300. .77280 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .82183 3900. .83145 4000. .84098 4100. .85045 4200. .85990 4300. .85910 4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
2000. .70027 2100. .69856 2200. .69856 2300. .70029 2400. .70345 2500. .70794 2600. .71358 2700. .72022 2800. .73584 3000. .74455 3100. .75368 3200. .76313 3300. .77280 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .8183 3900. .8145 4000. .86941 4400. .85990 4300. .88910 4600. .89959 4700. .89959 4700. .91079 4800. .92296 4900. .93640		
2100. .69856 2200. .69861 2300. .70029 2400. .70345 2500. .70794 2600. .71358 2700. .72022 2800. .72769 2900. .73584 3000. .74455 3100. .75368 3200. .76313 3300. .77280 3400. .78259 3500. .79245 3600. .81210 3800. .81145 4000. .84098 4100. .85045 4200. .85990 4300. .89590 4300. .89590 4300. .89590 4300. .89959 4300. .89959 4700. .91079 4800. .92296 4900. .93640		
2200. .69861 2300. .70029 2400. .70145 2500. .70794 2600. .71358 2700. .72022 2800. .72769 2900. .74455 3100. .74455 3200. .76313 3200. .76313 3300. .77280 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .82183 3900. .83145 4000. .84098 4100. .85045 4200. .85990 4300. .86941 4400. .87910 4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
2300. .70029 2400. .70345 2500. .70794 2600. .71358 2700. .72022 2800. .77354 3000. .74455 3100. .75368 3200. .76313 3300. .77280 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .82183 3900. .83145 4000. .84098 4100. .85045 4200. .85990 4300. .8959 4300. .8959 4300. .8959 4300. .8959 4500. .89959 4700. .91079 4800. .92296 4900. .93640		
2400. .70345 2500. .70794 2600. .71358 2700. .72022 2800. .72769 2900. .73584 3000. .74455 3100. .75368 3200. .76313 3300. .77280 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .8183 3900. .8145 4000. .84098 4100. .85045 4200. .85990 4300. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
2500. .70794 2600. .71358 2700. .72022 2800. .72769 2900. .73584 3000. .74455 3100. .75368 3200. .76313 3300. .77280 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .82183 3900. .83145 4000. .84098 4100. .85045 4200. .85990 4300. .86941 4400. .87910 4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
2700. .72022 2800. .72769 2900. .73584 3000. .74455 3100. .75368 3200. .75313 3300. .72280 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .8183 3900. .8145 4000. .84098 4100. .85045 4200. .85990 4300. .86941 4400. .87910 4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640	2500.	
2800. .72769 2900. .73584 3000. .7455 3100. .75368 3200. .76313 3300. .77280 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .82183 3900. .83145 4000. .86941 4100. .85990 4300. .86941 4400. .87910 4500. .8959 4700. .8959 4700. .91079 4800. .92296 4900. .93640	2600.	.71358
2900. .73584 3000. .74455 3100. .75368 3200. .76313 3300. .77280 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .82183 3900. .8145 4000. .84098 4100. .85045 4200. .85990 4300. .86941 4400. .87910 4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640	2700.	.72022
3000. .74455 3100. .75368 3200. .76313 3300. .77280 3400. .78259 3500. .78245 3600. .80230 3700. .81210 3800. .82183 3900. .83145 4000. .84098 4100. .85045 4200. .85990 4300. .86941 4400. .87910 4500. .88959 4300. .89959 4700. .91079 4800. .92296 4900. .93640	2800.	.72769
3100. .75368 3200. .76313 3300. .77280 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .82183 3900. .83145 4000. .86941 4100. .85990 4300. .86941 4400. .87910 4500. .8959 4700. .91079 4800. .92296 4900. .93640	2900.	.73584
3200. .76313 3300. .77280 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .82183 3900. .83145 4000. .84098 4100. .85045 4200. .85990 4300. .86941 4400. .87910 4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
3300. .77280 3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .82183 3900. .83145 4000. .84098 4100. .85045 4200. .85990 4300. .86941 4400. .87910 4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
3400. .78259 3500. .79245 3600. .80230 3700. .81210 3800. .82183 3900. .83145 4000. .84098 4100. .85990 4300. .85990 4300. .86941 4400. .87910 4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
3500. .79245 3600. .80230 3700. .81210 3800. .82183 3900. .83145 4000. .84098 4100. .85045 4200. .85990 4300. .86941 4400. .87910 4500. .88910 4600. .9959 4700. .91079 4800. .92296 4900. .93640		
3600. .80230 3700. .81210 3800. .82183 3900. .83145 4000. .84098 4100. .85045 4200. .85990 4300. .86941 4400. .87910 4500. .88910 4600. .9959 4700. .91079 4800. .92296 4900. .93640		
3700. .81210 3800. .82183 3900. .83145 4000. .84098 4100. .85045 4200. .85990 4300. .86941 4400. .87910 4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
3800. .82183 3900. .83145 4000. .84098 4100. .85045 4200. .85990 4300. .86941 4400. .87910 4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
3900. .83145 4000. .84098 4100. .85045 4200. .85990 4300. .86941 4400. .87910 4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
4000. .84098 4100. .85045 4200. .85990 4300. .86941 4400. .87910 4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
4100. .85045 4200. .85990 4300. .86941 4400. .87910 4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
4200. .85990 4300. .86941 4400. .87910 4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
4300. .86941 4400. .87910 4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
4400. .87910 4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
4500. .88910 4600. .89959 4700. .91079 4800. .92296 4900. .93640		
460089959 470091079 480092296 490093640		
470091079 480092296 490093640		
480092296 490093640		
490093640		
+JJL1/		
		= . =

Table B-12Compressibility Factor of Natural Gas

**************************************	SPECIFIC GRAVITY
80.	.650
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.95008
200.	.94295
300.	.93303
400.	.92081
500.	.90677
600.	.89136
700.	.87501
800.	.85809
900.	.84098
1000.	.82400
1100.	.80747
1200.	.79167
1300.	.77687
1400.	.76329
1500.	.75114
1600.	.74058
1700.	.73172
1800.	.72465
1900.	.71940
2000. 2100.	.71595
2200.	.71426 .71422
2300.	.71422
2400.	.71863
2500.	.72279
2600.	.72803
2700.	.73420
2800.	.74115
2900.	.74874
3000.	.75683
3100.	.76531
3200.	.77408
3300.	.78305
3400.	.79213
3500.	.80125
3600.	.81037
3700.	.81943
3800.	.82841
3900.	.83730
4000.	.84609
4100.	.85482
4200.	.86354
4300.	.87231
4400.	.88125
4500.	.89049
4600.	.90019
4700.	.91056
4800.	.92186
4900.	.93435
5000.	.94839

Table B-13Compressibility Factor of Natural Gas

Table B-14			
Compressibility Factor of Natural Ga	IS		

TEMPERATURE, OF 90.	SPECIFIC GRAVITY .650
50.	.030
	COMPRESSIBILITY FACTOR Z
100.	.95082
200.	.94455
300.	.93557
400.	.92435
500.	.91136
600.	.89701
700.	.88170
800.	.86582
900.	.84971
1000.	.83368
1100.	.81805
1200.	.80307
1300.	.78902
1400. 1500.	.77611
1600.	.76453 .75445
1700.	.74599
1800.	.73922
1900.	.73417
2000.	.73084
2100.	.72916
2200.	.72907
2300.	.73043
2400.	.73311
2500.	.73697
2600.	.74185
2700.	.74760
2800.	.75408
2900.	.76115
3000.	.76868
3100.	.77657
3200.	.78471
3300.	.79303
3400.	.80145
3500. 3600.	.80990
3700.	.81834 .82672
3800.	.83501
3900.	.83301
4000.	.85133
4000.	.85938
4200.	.86742
4300.	.87551
4400.	.88376
4500.	.89228
4600.	.90125
4700.	.91085
4800.	.92133
4900.	.93295
5000.	.94602

Table B-15			
Compressibility	Factor of	of Natural	Gas

	SPECIFIC GRAVITY
TEMPERATURE, OF 100.	.650
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.95149
200.	.94600
300.	.93790
400.	.92761
500.	.91559
600.	.90224
700.	.88793
800.	.87303
900.	.85786
1000.	.84274
1100.	.82796
1200.	.81378
1300.	.80044
1400.	.78817
1500.	.77715
1600.	.76755
1700.	.75947
1800.	.75300
1900.	.74816
2000.	.74495
2100.	.74331
2200.	.74317
2300.	.74441
2400.	.74691
2500.	.75051
2600.	.75506
2700.	.76044
2800.	.76648
2900.	.77307
3000.	.78009
3100.	.78743
3200.	.79501
3300.	.80273
3400.	.81054
3500.	.81837
3600.	.82618
3700.	.83393
3800.	.84160
3900.	.84917
4000.	.85665
4100.	.86408
4200.	.87149
4300.	.87894
4400.	.88655
4500.	.89442
4600.	.90270
4700.	.91159
4800.	.92131
4900.	.93211
5000.	.94429

Table B-16			
Compressibility Factor of Natural Gas	5		

TEMPERATURE, of	**************************************
60.	.700
	COMPRESSIBILITY FACTOR Z
100.	.93263
200.	.91942
300.	.90381
400.	.88617
500.	.86692
600.	.84646
700.	.82520
800.	.80352
900.	.78179
1000.	.76039
1100.	.73966
1200.	.71995
1300.	.70158
1400.	.68488
1500.	.67009
1600.	.65746
1700.	.64714
1800.	.63925
1900.	.63382
2000.	.63083
2100.	.63018
2200.	.63173
2300.	.63527
2400.	.64060 .64746
2500. 2600.	.65562
2800.	.65562
2800.	.67495
2900.	.68571
3000.	.69697
3100.	.70859
3200.	.72045
3300.	.73247
3400.	.74455
3500.	.75664
3600.	.76868
3700.	.78064
3800.	.79249
3900.	.80424
4000.	.81590
4100.	.82749
4200.	.83908
4300.	.85077
4400.	.86267
4500.	.87494
4600.	.88778
4700.	.90143
4800.	.91618
4900.	.93237
5000.	.95040

**************************************	SPECIFIC GRAVITY
70.	.700
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.93383
200.	.92188
300.	.90756
400.	.89125
500.	.87332
600.	.85417
700.	.83419
800.	.81374
900.	.79318
1000.	.77287
1100.	.75316
1200.	.73437
1300.	.71684
1400.	.70086
1500.	.68669
1600.	.67456
1700.	.66464
1800.	.65703
1900.	.65178
2000.	.64886 .64817
2100. 2200.	.64958
2300.	.65289
2400.	.65789
2500.	.66434
2600.	.67202
2700.	.68070
2800.	.69017
2900.	.70025
3000.	.71079
3100.	.72165
3200.	.73272
3300.	.74391
3400.	.75515
3500.	.76638
3600.	.77755
3700.	.78864
3800.	.79963
3900.	.81050
4000.	.82129
4100.	.83201
4200.	.84274
4300.	.85355
4400.	.86456
4500.	.87593
4600.	.88783
4700.	.90051
4800.	.91423
4900.	.92932
5000.	.94614

Та	ble B-1	7		
Compressibility	Factor	of I	Natural	Gas

Та	ble B-18		
Compressibility	Factor o	of Natural	Gas

TEMPERATURE, oF	SPECIFIC GRAVITY
80.	.700
	COMPRESSIBILITY FACTOR Z
100.	.93491
200.	.92411
300.	.91100
400.	.89592
500.	.87924
600.	.86133
700.	.84256
800.	.82329
900.	.80385
1000.	.78460
1100. 1200.	.76587
1300.	.74798
1400.	.73125 .71598
1400.	.70242
1600.	. 70242
1700.	.68128
1800.	.67397
1900.	.66891
2000.	.66607
2100.	.66538
2200.	.66668
2300.	.66980
2400.	.67451
2500.	.68061
2600.	.68785
2700.	.69603
2800.	.70494
2900.	.71440
3000.	.72428
3100.	.73444
3200.	.74477
3300. 3400.	.75521
3500.	.76567
3600.	.77611 .78649
3700.	.79677
3800.	.80695
3900.	.81702
4000.	.82700
4100.	.83693
4200.	.84685
4300.	.85684
4400.	.86704
4500.	.87756
4600.	.88860
4700.	.90037
4800.	.91313
4900.	.92718
5000.	.94288

TEMPERATURE, oF 90.	SPECIFIC GRAVITY .700
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.93589
200.	.92615
300.	.91416
400.	.90024
500.	.88473
600.	.86799
700.	.85037
800.	.83221
900.	.81385
1000.	.79561
1100.	.77782
1200.	.76080
1300.	.74485
1400.	.73027
1500.	.71731
1600.	.70619
1700.	.69708
1800.	.69007
1900.	.68522
2000.	.68250
2100.	.68182
2200.	.68304
2300.	.68600
2400.	.69047
2500.	.69625
2600.	.70310
2700.	.71083
2800.	.71923
2900.	.72814
3000.	.73741
3100.	.74693
3200.	.75659
3300.	.76633
3400.	.77608
3500.	.78579
3600.	.79543
3700.	.80498
3800.	.81441
3900.	.82374
4000.	.83298
4100.	.84215
4200.	.85133
4300.	.86058
4400.	.87001
4500.	.87975
4600.	.88998
4700.	.90091
4800.	.91277
4900.	.92586
5000.	.94051

Table B-19Compressibility Factor of Natural Gas

TEMPERATURE, OF 100.	SPECIFIC GRAVITY .700	
PRESSURE psia.	COMPRESSIBILITY FACTOR Z	
100.	.93677	
200.	.92801	
300.	.91706	
400.	.90422	
500.	.88981	
600.	.87418	
700.	.85765	
800.	.84055	
900.	.82321	
1000.	.80594	
1100.	.78906	
1200.	.77287	
1300.	.75769	
1400.	.74378	
1500.	.73141	
1600.	.72078	
1700.	.71207	
1800.	.70538	
1900.	.70074	
2000.	.69814	
2100.	.69750	
2200.	.69868	
2300.	.70150	
2400.	.70577	
2500.	.71127	
2600.	.71778	
2700. 2800.	.72510 .73304	
2900.	.73304	
3000.	.75016	
3100.	.75909	
3200.	.76814	
3300.	.77724	
3400.	.78633	
3500.	.79538	
3600.	.80434	
3700.	.81320	
3800.	.82195	
3900.	.83059	
4000.	.83914	
4000.	.84763	
4200.	.85612	
4200.	.85612	
4300.	.87340	
4400.	.88242	
4500.	.89190	
4800.	.90204	
4800.	.91307	
4800.	.92527	
5000.	.93894	

Table B-20Compressibility Factor of Natural Gas

Table B-21	
Compressibility Factor of Natural Ga	as

TEMPERATURE, oF 60.	SPECIFIC GRAVITY .750
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.91676
200.	.89894
300. 400.	.87920
400. 500.	.85783
600.	.83515 .81149
700.	.78720
800.	.76262
900.	.73811
1000.	.71403
1100.	.69075
1200.	.66866
1300.	.64813
1400.	.62953
1500.	.61318
1600.	.59936
1700.	.58829
1800.	.58009
1900.	.57483
2000.	.57244
2100.	.57280
2200.	.57572
2300.	.58092
2400.	.58813
2500.	.59701
2600.	.60728
2700.	.61862
2800.	.63080
2900.	.64357
3000.	.65675
3100.	.67019
3200.	.68378
3300.	.69742
3400.	.71106
3500.	.72463
3600.	.73811
3700.	.75147
3800.	.76471
3900.	.77782
4000.	.79084
4100.	.80381
4200.	.81680
4300.	.82990
4400.	.84324
4500.	.85699
4600.	.87136
4700.	.88660
4800.	.90302
4900.	.92099
5000.	.94090

TEMPERATURE, oF 70.	SPECIFIC GRAVITY .750
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.91829
200.	.90199
300.	.88377
400.	.86390
500.	.84268
600.	.82044
700.	.79751
800.	.77423
900.	.75093
1000.	.72797
1100.	.70573
1200.	.68457
1300.	.66487
1400.	.64698
1500.	.63124
1600.	.61791
1700.	.60721
1800.	.59927
1900.	.59415
2000.	.59179
2100.	.59208
2200.	.59481
2300.	.59974
2400.	.60657
2500.	.61499
2600.	.62472
2700.	.63545
2800.	.64694
2900.	.65898
3000.	.67137
3100.	.68398
3200.	.69671
3300.	.70946
3400.	.72217
3500.	.73482
3600.	.74735
3700.	.75977
3800.	.77206
3900.	.78423
4000.	.79630
4100.	.80832
4200.	.82036
4300.	.83250
4400.	.84487
4500.	.85764
4600.	.87099
4700.	.88517
4800.	.90047
4900.	.91723
5000.	.93584

Table B-22Compressibility Factor of Natural Gas

TEMPERATURE, oF 80.	SPECIFIC GRAVITY .750
PRESSURE psia.	COMPRESSIBILITY FACTOR 2
100.	.91968
200.	.90479
300.	.88798
400.	.86951
500.	.84968
600.	.82879
700.	.80716
800.	.78511
900.	.76299
1000.	.74113
1100.	.71989
1200.	.69965
1300.	.68076
1400.	.66358
1500.	.64844
1600.	.63561
1700.	.62530
1800.	.61764
1900.	.61268
2000.	.61039
2100.	.61064
2200.	.61323
2300.	.61792
2400.	.62442
2500.	.63244
2600.	.64168
2700.	.65186
2800.	.66274
2900.	.67410
3000.	.68577
3100.	.69763
3200.	.70955
3300.	.72148
3400.	.73336
3500.	.74514
3600.	.75681
3700.	.76835
3800.	.77976
3900.	.79105
4000.	.80225
4100.	.81339
4200.	.82454
4300.	.83580
4400.	.84727
4500.	.85911
4600.	.87151
4700.	.88470
4800.	.89895
4900.	.91458
5000.	.93197

Table B-23Compressibility Factor of Natural Gas

Compressibility Factor of Natural Gas		
**************************************	**************************************	
PRESSURE psia.	COMPRESSIBILITY FACTOR Z	
100		

Table B-24 Compressibility Factor of Natural Gas

TEMPERATURE, OF	SPECIFIC GRAVITY
90.	.750
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.92094
200.	.90735
300.	.89186
400.	.87470
500.	.85618
600.	.83657
700.	.81618
800.	.79532
900.	.77433
1000.	.75352
1100.	.73327
1200.	.71391
1300.	.69583
1400.	.67936
1500.	.66482
1600.	.65248
1700.	.64257
1800.	.63520
1900.	.63043
2000.	.62823
2100.	.62847
2200.	.63096
2300.	.63546
2400.	.64168
2500.	.64935
2600.	.65815
2700.	.66784
2800.	.67816
2900.	.68890
3000.	.69992
3100.	.71108
3200.	.72227
3300.	.73345
3400.	.74455
3500.	.75554
3600.	.76640
3700.	.77714
3800.	.78774
3900.	.79821
4000.	.80860
4100.	.81892
4200.	.82926
4300.	.83969
4400.	.85033
4500.	.86131
4600.	.87283
4700.	.88509
4800.	.89836
4900.	.91294
5000.	.92919

Table B-25		
Compressibility Factor of Natural Gas		

**************************************	SPECIFIC GRAVITY .750
PRESSURE psia.	COMPRESSIBILITY FACTOR Z
100.	.92209
200.	.90969
300.	.89543
400.	.87951
500.	.86221
600.	.84382
700.	.82461
800.	.80490
900.	.78499
1000.	.76520
1100.	.74589
1200.	.72741
1300.	.71011
1400.	.69432
1500.	.68038
1600.	.66855
1700.	.65904
1800.	.65198
1900.	.64742
2000.	.64532
2100.	.64558
2200.	.64800
2300.	.65234
2400.	.65833
2500.	.66568
2600.	.67411
2700.	.68335
2800.	.69316
2900.	.70336
3000.	.71378
3100.	.72430
3200.	.73483
3300.	.74531
3400.	.75569
3500.	.76595
3600.	.77608
3700.	.78607
3800.	.79592
3900.	.80565
4000.	.81528
4100.	.82485
4200.	.83443
4300.	.84410
4400.	.85396
4500.	.86415
4600.	.87485
4700.	.88625
4800.	.89861
4900.	.91221
5000.	.92739

Appendix C

COMPILING THE FORTRAN SOURCE CODE

The FORTRAN source code, like any high level code, is a compiler language, and the programmer must translate (compile) the code to be recognized by the computer. An intermediate program called a compiler is required to expedite this translation. The compiler used for translating (compiling) the programs in this book is the **Microsoft** (**MS**) **FORTRAN Version 5** compiler, developed and marketed by the Microsoft Corporation. Both FORTRAN Compiler Version 5 and the latest Version 5.1 are designed for compiling a FORTRAN source program on the personal computer. The MS-FORTRAN compiler consists of a set of floppy diskettes, a reference manual, and a user's guide. These give information about the files on the diskettes, and how they are used.

To compile the FORTRAN source code, the programmer needs a personal computer with a hard disk having a minimum of five megabyte disk space. The FORTRAN source program is written and edited by using any text editor or word processor. This is entered on the program with an appropriate filename, with the extension FOR. An object program (extension OBJ) and an executable program (extension EXE) with the same filename as the source program are created during compilation and stored on the program. The source code is created using the Microsoft FORTRAN text editor or a word processor in non-document mode. Using Microsoft text editor, the source code of a file PROG11.FOR can be created from the C:\> prompt, command line by typing:

C:\>m PROG11.FOR

The following procedure compiles and links the source code.

From the C:\>prompt command line, type

f1 /c /4Yb /FsCon PROG11.FOR

where /c = compiles without linking the source code /4Yb = Enables the extended error handling at run time

/FsCon = Produces a source code listing on the screen

The following messages will appear on the screen:

Microsoft ^(R) FORTRAN Optimizing Compiler Version 5.00

Copyright ^(c) Microsoft Corp 1982–1989. All rights reserved.

The link command invokes the Microsoft segmented executable linker.

link PROG11,,nul,llibfore/exe/nodef,nul

The following messages will appear on the screen:

Microsoft ^(R) Segmented - Executable Linker Version 5.03

Copyright ^(C) Microsoft Corp 1984–1989. All rights reserved.

A few seconds are required to compile the link step. When the drive light stops blinking, the C:\>prompt will return to the screen. Compilation is then complete.

A directory listing the files will include the following:

- PROG.FOR
- PROG11.OBJ
- PROG11.EXE
- PROG11.LST

Files with extensions MAP, and LST can be created and stored at the direction of the compiler program during compilation. The text for these files many be displayed, however only two files, PROG11.FOR and PROG11.LST will be displayed in readable form. The files with the extensions OBJ, MAP and EXE will be displayed in machine codes.

Executing the FORTRAN Program

Results of PROG11 are obtained by executing or running the program from the C:> prompt. The executable filename is typed either with or without the extension (EXE).

C:\>PROG11

After compilation has been completed, an executable file is created and stored, and the results are obtained. The files with extensions **FOR** and **EXE** are retained. Other files with the extensions **OBJ**, **MAP**, **LST**, and **BAK** resulting from editing the source program may be deleted to create more space on the hard disk drive.

Creating a Batch File

A batch file is a file containing one or more Disk Operating System (DOS) commands which are automatically executed one at a time when the batch command is initiated. A batch file can be assigned any suitable filename but must have the extension **BAT**. The batch file can be created by copying from the console (keyboard) with the internal COPY command or from a text editor or a word processor. Using the Microsoft text editor, creating a batch file **COMPILE.BAT** from compiling and running the computer program PROG11 involves the following sequence of commands.

From the C:\>prompt command line, type

C:>m COMPILE.BAT

The text editor screen is displayed and the following commands can then be typed.

@ECHO OFF @ECHO COMPILE THE SOURCE PROGRAM PROG11 f1 /c /4Yb /FsCon PROG11.FOR @ ECHO LINK THE OBJECT CODE OF PROGRAM PROG11 link PROG11,,,nul,llibfore/exe/nodef,nul @ECHO run the executable code of program PROG11 PROG11

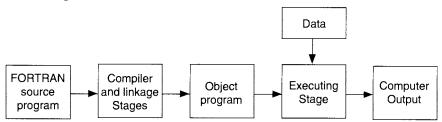
Errors

Errors detected during the compilation are located by the source program line number and appropriate numbered error messages are displayed. The error message numbers refer to the MS-FORTRAN error messages listed in the Microsoft FORTRAN User's Guide. Errors detected during the compilation are usually errors in FORTRAN syntax or procedure. Compilation is aborted when errors are detected, and the C:\>prompt is displayed. A source program may not be executed successfully even though no errors are detected during compilation. These are known as run time errors. These types of errors abort execution and display error messages to assist in identifying the errors. These error messages may be explicit and are sometimes difficult to detect. Examples of these errors are an attempt to calculate the square root of a negative number or divide by zero. A way of de-bugging errors is by having print or write statements at various stages of computation. Alternatively, a debugger software tool can be used to permit the programmer to execute his program one instruction at a time, and thus allows him to examine the state of his program variables after any instruction. Microsoft provides a high-level code degugger (CodeView) for this purpose.

Running the FORTRAN Program

Once the program has been typed into the computer using a text editor or a word processor in non-document mode and stored in a file, it must be compiled. A compiler is a program that translates a high-level language to machine language. The compilation step is the first stage in running the FORTRAN program on the computer. As the compiler translates statements, it also checks for syntax errors. These errors, also called compiler errors, are errors in the statements such as misspellings and punctuation errors. If these errors are detected, the compiler will print error messages or diagnostics for the programmer. After debugging the program, it can be recompiled using the MS-FORTRAN compiler. After a successful compilation, a linkage editor program performs the final stage before the program is executed.

The FORTRAN program is often referred to as the source program. The source code is converted into the machine language by the compiler and prepared for execution by the linkage editor. The linking process inserts pieces of the program in the right places in the object code to produce an executable program. This is sometimes known as a binary program, which is stored in a file. The data are inputted in the execution stage to obtain the output. This process is illustrated by the following block diagram.



The procedures given will enable users and readers of this book to readily compile and run all the source codes contained in the diskette.

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