Sterward. Prior to each acquisition, the rolled horseshoe nail market was highly concentrated and concentration increased substantially following each acquisition. The complaint alleges that entry into the production and sale of rolled horseshoe nails would be difficult and time consuming—taking well in excess of two years, entailing significant sunk costs, and requiring technical expertise.

The proposed Order would remedy the alleged violations by replacing the lost competition that has resulted from the acquisitions. The proposed Order would require Mustad to divest either (1) Capewell as an ongoing business, or (2) four fully functioning horseshoe nail machines, one spare nail machine, and grant a perpetual non-exclusive license to technology and know-how. In order to ensure that the acquirer of machinery would be able to quickly begin production at the same level of quality as exists currently, Mustad would be required to provide training and technical assistance to the acquirer for up to one year.

The proposed Order provides that Mustad shall divest Capewell or the machinery no later than May 15, 1996. If Mustad does not complete the required divestiture during the allotted time period, then a trustee may be appointed to divest the machinery within twelve months. The time period for the trustee to complete the divestiture may be extended twice.

The proposed Order requires Mustad to submit a report of compliance with the proposed Order's divestiture requirements within sixty (60) days following the date the proposed Order becomes final, and every sixty (60) days thereafter until Mustad has completed the divestiture.

Finally, the proposed Order prohibits Mustad from acquiring any interest in any other company engaged in, or attempting to engage in, the production or sale of horseshoe nails without giving prior notice to the Commission and observing certain waiting periods for a period of ten years.

The purpose of this analysis is to facilitate public comment on the proposed Order. This analysis is not intended to constitute an official interpretation of the Agreement or the proposed Order or in any way to modify the terms of the Agreement or the proposed Order.

#### Benjamin I. Berman,

Acting Secretary.

[FR Doc. 95-20142 Filed 8-14-95; 8:45 am]

BILLING CODE 6750-01-M

## **DEPARTMENT OF HEALTH AND HUMAN SERVICES**

## Centers for Disease Control and Prevention

**Clinical Laboratory Improvement** Advisory Committee (CLIAC) and Subcommittee on Proficiency Testing, **Quality Assurance, and Quality** Control; Meetings

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following Federal advisory committee meetings.

Name: Subcommittee on Proficiency Testing, Quality Assurance, and Quality Control, Clinical Laboratory Improvement Advisory Committee (CLIAC).

Time and Date: 8:30 a.m.-12 noon, August 30, 1995.

Place: Swissôtel Atlanta, 3391 Peachtree Road, NE, Atlanta, Georgia 30326.

Status: Open to the public, limited only by the space available.

Purpose: This subcommittee advises CLIAĈ on issues related to proficiency testing, quality assurance, and quality

Matters to be discussed: The Subcommittee will discuss quality control requirements for test method verification and appropriate materials for quality control testing.

Agenda items are subject to change as priorities dictate.

Name: Clinical Laboratory Improvement Advisory Committee.

Times and Dates: 1 p.m.-4:30 p.m., August 30, 1995., 8 a.m.-4 p.m., August 31, 1995. Place: Swissôtel Atlanta, 3391 Peachtree

Road, NE, Atlanta Georgia 30326 Status: Open to the public, limited only by

the space available.

Purpose: This committee is charged with providing scientific and technical advice and guidance to the Secretary of Health and Human Services, the Assistant Secretary for Health, and the Director, CDC, regarding the need for, and the nature of, revisions to the standards under which clinical laboratories are regulated; the impact of proposed revisions to the standards; and the modification of the standards to accommodate technological advances.

Matters to be discussed: The agenda will include an update on the implementation of the Clinical Laboratory Improvement Amendments (CLIA), a CDC presentation on CLIA quality control requirements, public presentations on quality control requirements, a discussion of the quality control requirements for the final regulations, and a summary of the meeting of the Subcommittee on Proficiency Testing, Quality Assurance, and Quality Control.

Agenda items are subject to change as priorities dictate.

Written comments on the quality control requirements are welcome. Comments should not exceed five single-spaced, typed pages in length and should be received by the contact person no later than August 24, 1995.

Anyone wishing to make an oral presentation that would include data pertinent to CLIA quality control requirements should submit their request, in writing, to the contact person by close of business, August 24, 1995. The request should include the name, address, and telephone number of the participant; the approximate time needed; and a brief summary of the topic and data to be presented. Depending on the number of requests, up to 10 minutes will be allowed for each oral presentation.

Contact Person for addition information: John C. Ridderhof, Dr. P.H., Division of Laboratory Systems, Public Health Practice Program Office, CDC, 4770 Buford Highway, NE, Mailstop G-25, Atlanta, Georgia 30341-3724, telephone 404/488-7660, FAX 404-488-7663

Dated: August 9, 1995.

#### Carolyn J. Russell,

Director, Management Analysis and Services Office Centers for Disease Control and Prevention (CDC).

[FR Doc. 95-20104 Filed 8-14-95: 8:45 am] BILLING CODE 4163-18-M

# **National Committee on Vital and Health** Statistics (NCVHS) Subcommittee on **Mental Health Statistics and NCVHS** Subcommittee on Disability and Long-**Term Care Statistics: Meetings**

Pursuant to Pub. L. 92–463, the National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC), announces the following subcommittee meetings.

Name: NCVHS Subcommittee on Mental Health Statistics and NCVHS Subcommittee on Disability and Long-Term Care Statistics.

Time and Date: 9 a.m.-5 p.m., September 12, 1995.

Place: Room 503A-529A, Hubert H. Humphrey Building, 200 Independence Avenue, SW, Washington, D.C. 20201.

Status: Open.

Purpose: The Subcommittee on Mental Health Statistics and the Subcommittee on Disability and Long-Term Care Statistics will meet jointly to consider and discuss presentations on a variety of payment and service models. There will be presentations on the Program of All-inclusive Care for the Elderly (PACE), the second phase of the Social Health Maintenance Organizations, consumer choice plans, and demonstration projects.

Name: NCVHS Subcommittee on Mental Health Statistics.

Time and Date: 9 a.m.-12 noon, September 13, 1995.

Place: Room 503A-529A, Hubert H. Humphrey Building, 200 Independence Avenue, SW, Washington, D.C. 20201. Status: Open.