Public telecommunications services means noncommercial educational and cultural radio and television programs, and related noncommercial instructional or informational material that may be transmitted by means of electronic communications.

* * * * *

3. Section 2301.4(a) is revised to read as follows:

§ 2301.4 Eligible organizations and scope of projects.

(a) Eligible applicants (Construction and Planning Grants). In order to apply for and receive a PTFP Construction or Planning Grant, an applicant must be:

(1) A public or noncommercial educational broadcast station;

(2) A noncommercial telecommunications entity;

(3) A system of public telecommunications entities;

(4) A nonprofit foundation, corporation, institution, or association organized primarily for educational or cultural purposes; or

(5) A state or local government (or agency thereof), or a political or special purpose subdivision of a state.

* * * * *

4. Section 2301.4 is further amended by removing paragraph (b), redesignating paragraph (c) as (b), and revising the newly designated paragraph (b) to read as follows:

* * * * *

(b) Scope of projects. An applicant that is eligible under paragraph (a) of this section may file an application with the agency for a planning or construction grant to achieve the following:

(1) The provision of new public telecommunications facilities to extend service to areas currently not receiving public telecommunications services;

(2) The expansion of the service areas of existing public telecommunications entities;

(3) The establishment of new public telecommunications entities serving areas currently receiving public telecommunications services; or

(4) The improvement of the capabilities of existing licensed public broadcast stations to provide public telecommunications services.

* * * * *

5. Paragraphs (d) through (f) of § 2301.4 are redesignated paragraphs (c) through (e) respectively.

[FR Doc. 95–31089 Filed 12–21–95; 8:45 am]

BILLING CODE 3510–60–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 165

[Docket No. 88P–0030]

RIN 0910–AA11

Beverages: Bottled Water; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a final rule that appeared in the Federal Register of November 13, 1995 (60 FR 57076). The document established a standard of identity for bottled water; recodified the standard of quality for bottled water; revised the definition of bottled water to include mineral water and ingredient uses of this product; and defined ''artesian water,'' ''ground water,'' ''mineral water,'' ''purified water,'' ''sparkling bottled water,'' ''spring water,'' ''sterile water,'' and ''well water.'' The document was published with some errors. This document corrects those errors.

EFFECTIVE DATE: May 13, 1996.

FOR FURTHER INFORMATION CONTACT: Shellee A. Davis, Center for Food Safety and Applied Nutrition (HFS–306), Food and Drug Administration, 200 C St. SW., Washington, DC 20204,202–205–4681.

In FR Doc. 95–27798, appearing on page 57076 in the Federal Register of Monday, November 13, 1995, the following corrections are made in § 165.110 *Bottled water*:

§ 165.110 [Corrected]

1. On page 57124, in the third column, in paragraph (a)(2)(v), in the fourth line, the phrase ''amount of carbon dioxide that'' is corrected to read ''amount of carbon dioxide from the source that''.

2. On page 57126, in the first column, in paragraph (b)(4)(i)(C)(*4*), beginning in the third line, ''Method 501.2'' is removed.

Dated: December 15, 1995.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 95–31200 Filed 12–21–95; 8:45 am]

BILLING CODE 4160–01–F

21 CFR Part 558

New Animal Drugs For Use In Animal Feeds; Zoalene

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect approval of a supplemental new animal drug application (NADA) filed by A. L. Pharma, Inc. The supplemental NADA provides for wider assay limits for zoalene Type A medicated articles. FDA is amending the regulation concerning medicated feed applications to reflect the approval.

EFFECTIVE DATE: December 22, 1995.

FOR FURTHER INFORMATION CONTACT: William G. Marnane, Center for Veterinary Medicine (HFV–143), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–2701.

SUPPLEMENTARY INFORMATION: A. L. Pharma, Inc., One Executive Dr., P.O. Box 1399, Fort Lee, NJ 07024, filed supplemental NADA 11–116, which provides for widening the current assay limits for zoalene Type A medicated articles. FDA reviewed the data and information submitted and is amending 21 CFR 558.4 to provide for an assay range of 92 to 104 percent for zoalene Type A medicated articles.

List of Subjects in 21 CFR Part 558

Animal drugs, Animal feeds.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR part 558 is amended as follows:

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

1. The authority citation for 21 CFR part 558 continues to read as follows:

Authority: Secs. 512, 701 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b, 371).

§ 558.4 [Amended]

2. Section 558.4 *Medicated feed applications* is amended in paragraph (d) in the ''Category I'' table in the entry for ''Zoalene'' in the second column by removing ''98–104'' and adding in its place ''92–104''.