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VIII. Effective Date and Congressional Notification

This Final Rule will take effect June 24, 2002. The Commission has determined, with the concurrence of the Administrator of the Office of Information and Regulatory Affairs of the Office of Management and Budget, that this rule is not a "major rule" within the meaning of Section 251 of the Small Business Regulatory Enforcement Fairness Act of 1996.⁷ The Commission will submit the Final Rule to both houses of Congress and the General Accounting Office.⁸

List of Subjects

18 CFR Part 35

Electric power rates, Electric utilities, Reporting and recordkeeping requirements.

18 CFR Part 141

Electric power, Reporting and recordkeeping requirements.

18 CFR Part 385

Administrative practice and procedure, Electric power, Penalties, Pipelines, Reporting and recordkeeping requirements.

By the Commission.

Magalie R. Salas,
Secretary.

In consideration of the foregoing, the Commission amends parts 35, 141 and 385, Chapter I, Title 18, of the Code of Federal Regulations, as follows:

PART 35—FILING OF RATE SCHEDULES

1. The authority citation for part 35 continues to read as follows:

Authority: 16 U.S.C. 791a–825r, 2601–2645; 31 U.S.C. 9701; 42 U.S.C. 7101–7352.

§ 35.25 [Amended]

2. In § 35.25, paragraph (c)(1)(ii)(C) is removed, and paragraph (c)(1)(ii)(D) is redesignated as paragraph (c)(1)(ii)(C).

PART 141—STATEMENTS AND REPORTS (SCHEDULES)

3. The authority citation for part 141 continues to read as follows:

Authority: 15 U.S.C. 79; 16 U.S.C. 791a–828c, 2601–2645; 31 U.S.C. 9701; 42 U.S.C. 7101–7352.

4. In § 141.1, paragraphs (b)(1)(i) and (b)(2) are revised to read as follows:

§ 141.1 FERC No. Form 1, Annual report of Major electric utilities, licensees and others.

* * * * *

(b) *Filing requirements.* (1) *Who must file—(i) Generally.* Each Major electric utility (as defined in part 101 of Subchapter C of this chapter) and other entity, *i.e.* each corporation, person or licensee as defined in section 3 of the Federal Power Act (16 U.S.C. 792 *et seq.*), including any agency, authority or other legal entity or instrumentality engaged in generation, transmission, distribution, or sale of electric energy, however produced, throughout the United States and its possessions, having sales or transmission service equal to Major as defined above, whether or not the jurisdiction of the Commission is otherwise involved, shall prepare and file electronically with the Commission the FERC Form 1 pursuant to the General Instructions set out in that form.

* * * * *

(2) *When to file and what to file.* This report shall be filed on or before April 30 of each year for the previous calendar year. This report must be filed with the Federal Energy Regulatory Commission as prescribed in § 385.2011 of this chapter and as indicated in the General Instructions set out in this form, and must be properly completed and verified. Filing on electronic media pursuant to § 385.2011 of this chapter will be required commencing with the report required to be submitted for the reporting calendar year of 2002, due on or before April 30, 2003.

PART 385—RULES OF PRACTICE AND PROCEDURE

5. The authority citation for part 385 continues to read as follows:

Authority: 5 U.S.C. 551–557; 15 U.S.C. 717–717z, 3301–3432; 16 U.S.C. 791a–825r, 2601–2645; 31 U.S.C. 9701; 42 U.S.C. 7101–7352; 49 U.S.C. 60502; 49 App. U.S.C. 1–85 (1988).

6. In § 385.2011, paragraph (c)(3) is revised to read as follows:

§ 385.2011 Procedures for filing on electronic media (Rule 2011).

* * * * *

(c) *What to file.* * * *

(3) With the exception of the Form 1, the electronic media must be accompanied by the traditional prescribed number of paper copies.

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[FR Doc. 02–12798 Filed 5–22–02; 8:45 am]

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DEPARTMENT OF THE TREASURY

Customs Service

19 CFR Part 141

[T.D. 02–28]

Technical Amendment to the Customs Regulations: Reusable Shipping Devices Arriving From Canada and Mexico

AGENCY: Customs Service, Department of the Treasury.

ACTION: Final rule.

SUMMARY: This document amends the Customs Regulations to include certain reusable shipping devices arriving from Canada or Mexico in the list of merchandise excepted from the requirement that all merchandise imported into the United States be entered. The substantive regulation allowing for these types of devices to be excepted from entry is set forth in § 10.41b(b) of the Customs Regulations. During a periodic review of its regulations to ensure that they are current, correct and consistent, Customs noted that in § 141.4 of the Customs Regulations, the list of merchandise excepted from the entry requirement did not cross-reference § 10.41b(b). This document remedies that omission.

EFFECTIVE DATE: May 23, 2002.

FOR FURTHER INFORMATION CONTACT: Glen Vereb, Office of Regulations and Rulings, 202–927–1327.

SUPPLEMENTARY INFORMATION:

⁷ 5 U.S.C. 804(2).

⁸ 5 U.S.C. 801(a)(1)(A).

Background

Under § 141.4(a), Customs Regulations (19 CFR 141.4(a)), all merchandise imported into the United States is required to be entered, unless specifically excepted. The exceptions from the general rule that all imported merchandise must be entered are set forth in § 141.4(b), Customs Regulations (19 CFR 141.4(b)). In particular, § 141.4(b)(3) excepts instruments of international trade as described in § 10.41a, Customs Regulations (19 CFR 10.41a). In addition to this exemption from entry, however, certain reusable shipping devices arriving from Canada or Mexico are also exempted from entry pursuant to § 10.41b(b), Customs Regulations (19 CFR 10.41b(b)), as amended by Treasury Decision (T.D.) 96-20 (61 FR 7987) of March 1, 1996.

Accordingly, this document amends § 141.4(b)(3), Customs Regulations (19 CFR 141.4(b)(3)), to include a reference to § 10.41b(b), in order to reflect that reusable shipping devices from Canada or Mexico are also exempted from Customs entry requirements. Furthermore, a reference is added in § 141.4(b)(3) to Chapter 98, Subchapter III, U.S. Note 3, Harmonized Tariff Schedule of the United States (HTSUS), which provides the underlying legal authority for the exemption of the specified shipping devices from Customs entry requirements.

Administrative Procedure Act, the Regulatory Flexibility Act and Executive Order 12866

Because the amendments merely conform with existing law or regulation, notice and public procedure are unnecessary, and for the same reason, pursuant to 5 U.S.C. 553(d)(3), a delayed effective date is not required. Because no notice of proposed rulemaking is required, the provisions of the Regulatory Flexibility Act (5 U.S.C. 601 *et seq.*) do not apply. Nor do these amendments meet the criteria for a "significant regulatory action" as specified in Executive Order 12866.

Drafting Information

The principal author of this document was Janet L. Johnson, Regulations Branch, Office of Regulations and Rulings, U.S. Customs Service. However, personnel from other offices participated in its development.

List of Subjects In 19 CFR Part 141

Customs duties and inspection, Entry of merchandise, Release of merchandise, Reporting and recordkeeping requirements.

Amendment to the Regulation

Part 141, Customs Regulations (19 CFR 141) is amended as set forth below.

PART 141—ENTRY OF MERCHANDISE

1. The general authority citation for Part 141, Customs Regulations, continues to read, and the specific sectional authority for § 141.4 is revised to read, as follows:

Authority: 19 U.S.C. 66, 1448, 1484, 1624.

Section 141.4 also issued under 19 U.S.C 1202 (General Note 19; Chapter 86, Additional U.S. Note 1; Chapter 89, Additional U.S. Note 1; Chapter 98, Subchapter III, U.S. Notes 3 and 4; Harmonized Tariff Schedule of the United States), 1498;

2. Section 141.4 is amended by revising paragraph (b)(3) to read as follows:

§ 141.4 Entry required.

(b) *Exceptions.*

(3) Instruments of international traffic described in § 10.41a and § 10.41b(b) of this chapter, under the conditions provided for in those sections. See also Chapter 98, Subpart III, U.S. Notes 3 and 4, HTSUS.

Robert C. Bonner,
Commissioner of Customs.

Approved: May 17, 2002.

Timothy E. Skud,
Deputy Assistant Secretary of the Treasury.
[FR Doc. 02-12938 Filed 5-22-02; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

New Animal Drugs for Use in Animal Feeds; Decoquinat and Chlortetracycline

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 new animal drug application (NADA) filed by Alpharma, Inc. The NADA provides for use of approved decoquinat and

chlortetracycline Type A medicated articles to make two-way combination Type B and Type C medicated feeds for calves, beef, and nonlactating dairy cattle used for prevention of coccidiosis, treatment of bacterial enteritis, and treatment of bacterial pneumonia.

DATES: This rule is effective May 23, 2002.

FOR FURTHER INFORMATION CONTACT:

Janis R. Messenheimer, Center for Veterinary Medicine (HFV-135), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301-827-7578, e-mail: [jmessen@cvm.fda.gov](mailto:jmessenh@cvm.fda.gov).

SUPPLEMENTARY INFORMATION: Alpharma, Inc., One Executive Dr., P.O. Box 1399, Fort Lee, NJ 07024, filed NADA 141-185 that provides for use of DECCOX (decoquinat) and AUREOMYCIN (chlortetracycline) Type A medicated articles to make combination drug Type B and Type C medicated feeds for calves, beef and nonlactating dairy cattle. The combination Type C feeds are used for the prevention of coccidiosis caused by *Eimeria bovis* and *E. zuernii*, for treatment of bacterial enteritis caused by *Escherichia coli*, and for treatment of bacterial pneumonia caused by *Pasteurella multocida* organisms susceptible to chlortetracycline. The NADA is approved as of March 15, 2002, and the regulations are amended in 21 CFR 558.195 to reflect the approval. The basis of approval is discussed in the freedom of information summary.

In accordance with the freedom of information provisions of 21 CFR part 20 and 514.11(e)(2)(ii), a summary of safety and effectiveness data and information submitted to support approval of each application may be seen in the Dockets Management Branch (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, between 9 a.m. and 4 p.m., Monday through Friday.

FDA has determined under 21 CFR 25.33(a)(2) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

This rule does not meet the definition of "rule" in 5 U.S.C. 804(3)(A) because it is a rule of "particular applicability." Therefore, it is not subject to the congressional review requirements in 5 U.S.C. 801-808.

List of Subjects in 21 CFR Part 558

Animal drugs, Animal feeds.