corrects that amendment number. In all other respects, the original document remains the same.

DATES: Effective October 25, 2004.

FOR FURTHER INFORMATION CONTACT: Tomaso DiPaolo, Aerospace Engineer, Chicago Aircraft Certification Office, FAA, Small Airplane Directorate, 2300 East Devon Avenue, Des Plaines, IL 60018; telephone: (847) 294–7031; fax: (847) 294–7834.

SUPPLEMENTARY INFORMATION: A final rule; request for comments AD, FR Doc. 04–22728, that applies to certain Hartzell Propeller Inc. (formerly Hartzell Propeller Products Division) Model HC–B5MP–3( )/M10282A( )+6 five bladed propellers, was published in the **Federal Register** on October 14, 2004, (69 FR 60952). The following correction is needed:

§ 39.13 [Corrected]

On page 60953, in the third column, in the Amendatory Language, in the first paragraph, in the fifth line, ''Amendment 39–XXXXX'' is corrected to read ''Amendment 39–13822''.

Issued in Burlington, MA, on October 18, 2004.

Francis A. Favara,

Acting Manager, Engine and Propeller Directorate, Aircraft Certification Service.

[FR Doc. 04–23815 Filed 10–22–04; 8:45 am]

BILLING CODE 4910–13–P

FEDERAL TRADE COMMISSION

16 CFR Part 305

Rule Concerning Disclosures Regarding Energy Consumption and Water Use of Certain Home Appliances and Other Products Required Under the Energy Policy and Conservation Act (''Appliance Labeling Rule'')

AGENCY: Federal Trade Commission.

ACTION: Final rule.

SUMMARY: The Federal Trade Commission (''Commission'') announces that the current ranges of comparability for refrigerators, refrigerator-freezers, and freezers will remain in effect until further notice.

DATES: Effective January 24, 2004.

FOR FURTHER INFORMATION CONTACT: Hampton Newsome, Attorney, Division of Enforcement, Federal Trade Commission, Washington, DC 20580 (202–326–2889); *hnewsome@ftc.gov.*

SUPPLEMENTARY INFORMATION: The Appliance Labeling Rule (''Rule'') was issued by the Commission in 1979, 44 FR 66466 (November 19, 1979), in response to a directive in the Energy Policy and Conservation Act of 1975 (''EPCA'').[1] The Rule covers several categories of major household appliances including refrigerators, refrigerator-freezers, and freezers.

I. Background

The Rule requires manufacturers of all covered appliances to disclose specific energy consumption or efficiency information (derived from the DOE test procedures) at the point of sale in the form of an ''EnergyGuide'' label, fact sheets (for some appliances), and in catalogs. The Rule requires manufacturers to include, on labels and fact sheets, an energy consumption or efficiency figure and a ''range of comparability.'' This range shows the highest and lowest energy consumption or efficiencies for all comparable appliance models so consumers can compare the energy consumption or efficiency of other models similar to the labeled model. The Rule also requires manufacturers to include, on labels for some products, including those that are the subject of this notice, a secondary energy usage disclosure in the form of an estimated annual operating cost based on a specified DOE national average cost for the fuel the appliance uses.

Section 305.8(b) of the Rule requires manufacturers, after filing an initial report, to report certain information annually to the Commission by specified dates for each product type.[2] These reports, which are to assist the Commission in preparing the ranges of comparability, contain the estimated annual energy consumption or energy efficiency ratings for the appliances derived from tests performed pursuant to the DOE test procedures. Because manufacturers regularly add new models to their lines, improve existing models, and drop others, the data base from which the ranges of comparability are calculated is constantly changing. To keep the required information on labels consistent with these changes, the Commission will publish new ranges if an analysis of the new information indicates that the upper or lower limits of the ranges have changed by more than 15%. Otherwise, the Commission will publish a statement that the prior ranges remain in effect for the next year.

II. 2004 Refrigerator Data

The annual submissions of data for refrigerators, refrigerator-freezers, and freezers have been made and analyzed by the Commission. The ranges of comparability for the products have not changed significantly for these products.[3] Therefore, the current ranges for these products (16 CFR Part 305, Appendices A1 through A8 and B1 through B3) will remain in effect until further notice.[4]

List of Subjects in 16 CFR Part 305

Advertising, Energy conservation, Household appliances, Labeling, Reporting and recordkeeping requirements.

■ The authority citation for part 305 continues to read as follows:

Authority: 42 U.S.C. 6294.

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 04–23820 Filed 10–24–04; 8:45 am]

BILLING CODE 6750–01–M

[1] 42 U.S.C. 6294. The statute also requires the Department of Energy (''DOE'') to develop test procedures that measure how much energy the appliances use, and to determine the representative average cost a consumer pays for the different types of energy available.

[2] Reports for refrigerators, refrigerator-freezers, and freezers are due August 1.

[3] The Commission's analysis excluded models with energy consumption figures that do not meet the current DOE energy conservation standards. *See* 62 FR 23102 (April 28, 1997).

[4] *See* November 19, 2001 (66 FR 57867), November 26, 2001 (66 FR 59050), December 10, 2001 (66 FR 63749), January 29, 2002 (67 FR 4173), and November 21, 2003 (68 FR 65631).

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 520

Oral Dosage Form New Animal Drugs; Praziquantel Tablets

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 abbreviated new animal drug application (ANADA) filed by Phoenix Scientific, Inc. The supplemental ANADA provides for use of oral praziquantel tablets for the removal of certain tapeworm parasites in dogs.

DATES: This rule is effective October 25, 2004.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV–104), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 301–827–8549, e-mail: *lonnie.luther@fda.gov*.

SUPPLEMENTARY INFORMATION: Phoenix Scientific, Inc., 3915 South 48th St. Ter., St. Joseph, MO 64503, filed a supplement to ANADA 200–265 that provides for use of PRAZI–C (praziquantel) Tablets for the removal of certain tapeworm parasites in dogs. Phoenix Scientific, Inc.'s PRAZI–C Tablets are approved as a generic copy of Bayer HealthCare LLC's Tape Worm Tabs approved under NADA 111–798. The supplemental ANADA is approved as of September 15, 2004, and the regulations are amended in 21 CFR 520.1870 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 21 CFR 514.11(e)(2)(ii), a summary of safety and effectiveness data and information submitted to support approval of this application may be seen in the Division of Dockets Management (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.

The agency has determined under 21 CFR 25.33(a)(1) 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 520

Animal drugs.

■ 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 520 is amended as follows:

PART 520—ORAL DOSAGE FORM NEW ANIMAL DRUGS

■ 1. The authority citation for 21 CFR part 520 continues to read as follows:

Authority: 21 U.S.C. 360b.

■ 2. Section 520.1870 is amended by revising paragraph (b)(2) to read as follows:

§ 520.1870 Praziquantel tablets.

* * * * *

(b) * * *

(2) No. 059130 for use of the product described in paragraph (a)(1) of this section, as in paragraph (c)(1) of this section.

* * * * *

Dated: October 14, 2004.

Steven D. Vaughn,

Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.

[FR Doc. 04–23761 Filed 10–22–04; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 524

Ophthalmic and Topical Dosage Form New Animal Drugs; Ivermectin Topical Solution

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 an abbreviated new animal drug application (ANADA) filed by Norbrook Laboratories, Ltd. The ANADA provides for topical use of ivermectin on cattle for treatment and control of various species of external and internal parasites.

DATES: This rule is effective October 25, 2004.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV–104), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 301–827–8549, e-mail: *lonnie.luther@fda.gov*.

SUPPLEMENTARY INFORMATION: Norbrook Laboratories, Ltd., Station Works, Newry BT35 6JP, Northern Ireland, filed ANADA 200–272 for NOROMECTIN (ivermectin) Pour On for Cattle. The application provides for topical use of 0.5 percent ivermectin solution on cattle for the treatment and control of various species of gastrointestinal nematodes, lungworms, grubs, horn flies, lice, and mites. Norbrook Laboratories, Ltd.'s NOROMECTIN Pour-On for Cattle is approved as a generic copy of Merial Ltd.'s IVOMEC Pour-On for Cattle, approved under NADA 140–841. The application is approved as of September 13, 2004, and the regulations are amended in 21 CFR 524.1193 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 21 CFR 514.11(e)(2)(ii), a summary of safety and effectiveness data and information submitted to support approval of this application may be seen in the Division of Dockets Management (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.

The agency has determined under 21 CFR 25.33(a)(1) 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 524

Animal drugs.

■ 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 524 is amended as follows:

PART 524—OPHTHALMIC AND TOPICAL DOSAGE FORM NEW ANIMAL DRUGS

■ 1. The authority citation for 21 CFR part 524 continues to read as follows:

Authority: 21 U.S.C. 360b.

§ 524.1193 [Amended]

■ 2. Section 524.1193 is amended in paragraph (b)(2) by adding in numerical order ''055529''.

Dated: October 8, 2004.

Steven D. Vaughn,

Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.

[FR Doc. 04–23760 Filed 10–22–04; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF THE TREASURY

Internal Revenue Service

26 CFR Parts 1 and 602

[TD 9160]

RIN 1545–AY35

Information Reporting Under Section 6050P for Discharges of Indebtedness

AGENCY: Internal Revenue Service (IRS), Treasury.

ACTION: Final regulations.