FDC date	State	City	Airport	FDC No.	Subject
04/16/02	TX	GREENVILLE	MAJORS	2/3085	NDB OR GPS RWY 17, AMDT 5B
04/16/02	TX	GREENVILLE	MAJORS	2/3086	TACAN RWY 17, AMDT 2A ROW
04/17/02	IL	CHICAGO/AURORA	AURORA MUNI	2/3099	VOR RWY 15, ORIG–A
04/17/02	TN	DICKSON	DICKSON MUNI	2/3126	VOR/DME OR GPS RWY 17, AMDT 4
04/17/02	TN	DICKSON	DICKSON MUNI	2/3127	NDB RWY 17, AMDT 2
04/17/02	NV	LAS VEGAS	McCARRAN INTL	2/3131	ILS RWY 25L, AMDT 3
04/18/02	TX	DALLAS-FORTH WORTH.	DALLAS-FORT WORTH INTERNATIONAL.	2/3171	RNAV (GPS) RWY 13R, ORIG
04/18/02	TX	DALLAS-FORT WORTH	DALLAS-FORT WORTH INTERNATIONAL.	2/3172	ILS RWY 13R, AMDT 6
04/18/02	TX	McKINNEY	McKINNEY MUNI	2/3178	GPS RWY 17, ORIG-B
04/18/02	TX	ATLANTA	HALL-MILLER	2/3179	RNAV (GPS) RWY 5, ORIG
04/18/02	VA	RICHMOND/ASHLAND	HANOVER COUNTY MUNI	2/3184	NDB RWY 16, ORIG-C
04/18/02	CA	LOS ANGELES	LOS ANGELES INTL	2/3204	ILS RWY 24R (CAT I, II, III) AMDT 22
04/19/02	VA	ROANOKE	ROANOKE REGIONAL/WOODRUM	2/3228	LDA RWY 6, AMDT 7B
04/19/02	KS	WICHITA	CESSNA AIRCRAFT FIELD	2/3256	VOR OR GPS-C, ORIG-A
04/19/02	WV	PINEVILLE	KEE FIELD	2//3258	GPS RWY 7, ORIG
04/19/02	WV	PINEVILLE	KEE FIELD	2/3259	GPS RWY 25, ORIG
04/19/02	VA	ROANOKE	ROANOKE REGIONAL/WOODRUM	2/3262	ILS RWY 33, AMDT 11
04/19/02	NY	ROCHESTER	GREATER ROCHESTER INTL	2/3280	ILS RWY 4, AMDT 17
04/19/02	NY	ROCHESTER	GREATER ROCHESTER INTL	2/3281	ILS RWY 22, AMDT 5. THIS REPLACES FDC 2/2747 IN TL02–11.
04/19/02	NY	ROCHESTER	GREATER ROCHESTER INTL	2/3283	RNAV (GPS) RWY 22, ORIG-A. THIS REPLACES FDC 2/2752 IN TLOS–11.
04/19/02	AK	ANCHORAGE	TED STEVENS ANCHORAGE INTL	2/3284	NDB RWY 6R, AMDT 6E
04/19/02	NY	ROCHESTER	GREATER ROCHESTER INTL	2/3286	ILS RWY 4, (CAT II), AMDT 17. THIS REPLACES FDC 2/2746 IN TL02–11.
04/22/02	SC	UNION	UNION COUNTY-TROY SHELTON FIELD.	2/3348	NDB RWY 5, ORIG
04/22/02	CA	JACKSON	WESTOVER FIELD AMADOR COUNTY.	2/3364	GPS RWY 1, ORIG
04/22/02	CA	JACKSON	WESTOVER FIELD AMADOR COUNTY.	2/3365	VOR/DME RWY 1, AMDT 1
04/22/02	CA	SACRAMENTO	McCLELLAN AIRFIELD	2/3367	ILS RWY 16, ORIG-A
04/23/02	VA	RICHMOND/ASHLAND	HANOVER COUNTY MUNI	2/3383	GPS RWY 16, AMDT 1A

[FR Doc. 02–10940 Filed 5–1–02; 8:45 am]
BILLING CODE 4910–13–M

DEPARTMENT OF ENERGY

Federal Energy Regulatory Commission

18 CFR Part 388

[Docket No. RM02–8–000; Order No. 625]

Revised Fees for Record Requests; Final Rule

Issued April 26, 2002.

AGENCY: Federal Energy Regulatory Commission.

ACTION: Final rule.

SUMMARY: The Federal Energy Regulatory Commission is amending its regulations to increase the fee for hard copies of documents printed from the Federal Energy Regulatory Records Information System (FERRIS) from 15 to 20 cents per page. This change is necessary due to decreased volume and will enable the Commission to continue offering copying services to the public.

EFFECTIVE DATE: This final rule is effective immediately upon issuance.

FOR FURTHER INFORMATION CONTACT: Katherina Quijada-Cusack, Office of the Chief Information Officer, Federal Energy Regulatory Commission, 888 First Street, NE., Washington, DC 20426, (202) 208–1748, *Katherina.Quijada-Cusack@ferc.gov.*

SUPPLEMENTARY INFORMATION:

I. Introduction

The Federal Energy Regulatory Commission is amending Section 388.109 of its regulations to increase the fee for hard copies of documents available through its Public Reference Room in electronic form from 15 to 20 cents per page.

II. Background

The Commission makes public documents available for download through the Internet.[1] Until recently, this has been done primarily through the Commission's Records and Information Management System (RIMS). The Commission now is in the process of replacing RIMS and other records systems with the Federal Energy Regulatory Records Information System (FERRIS). FERRIS will provide improved functionality and reliability to members of the public seeking information about Commission proceedings and other matters.[2]

Documents available electronically are also available to the public in hard copy. Currently, the Commission's regulations call for a charge of 15 cents per page for hard copies of documents that are available in electronic format.[3]

[1] 18 CFR 388.106.

[2] *See* 67 FR 10910 (Mar. 11, 2002).

[3] 18 CFR 388.109(a)(4)(i).

This rule will change the charge to 20 cents per page.

III. Discussion

Due to increased usage of the Internet by members of the public who wish to access public Commission documents, the Commission has seen a decreased demand for hard copies of electronically available documents. Because of the smaller volume, the Commission's Public Reference Room contractor, which was recently selected through a competed procurement as offering the best value among available firms, requires an increase in the copying charge for the service to continue to remain economically viable. Commission staff monitors printing statistics and has verified the contractor's need. The Commission does not believe the price increase will cause any hardship, particularly given the increasing reliance on electronic means for accessing documents. This final rule also deletes the reference to the Commission's Records and Information Management System (RIMS) and substitutes a reference to the new Federal Energy Regulatory Records Information System (FERRIS).

IV. Regulatory Flexibility Act Certification

The Regulatory Flexibility Act (''RFA'') requires agencies to prepare certain statements, descriptions, and analyses of proposed rules that will have a significant economic impact on a substantial number of small entities.[4] The Commission is not required to make such an analysis if a rule would not have such an effect.

The Commission does not believe that this rule would have such an impact on small entities. Charges for hard copies of documents remain modest and the Commission considers it very unlikely that any person or entity would require such a large volume of documents for this increase to have a significant impact.

V. Environmental Statement

Issuance of this Final Rule does not represent a major federal action having a significant adverse effect on the human environment under the Commission's regulations implementing the National Environmental Policy Act.[5] Part 380 of the Commission's regulations lists a number of exemptions where an Environmental Analysis or Environmental Impact Statement will not be done. Included are exemptions for procedural, ministerial or internal administrative actions, and for information gathering, analysis and dissemination.[6] This rulemaking is exempt under those provisions.

VI. Information Collection Statement

The Office of Management and Budget's (''OMB's'') regulations require that OMB approve certain information collection requirements imposed by agency rule.[7] This Final Rule contains no information reporting requirements, and is not subject to OMB approval.

VII. Document Availability

In addition to publishing the full text of this document in the **Federal Register**, the Commission provides all interested persons an opportunity to view and/or print the contents of this document via the Internet through FERC's Home Page (*http://www.ferc.gov*) and in FERC's Public Reference Room during normal business hours (8:30 a.m. to 5:00 p.m. Eastern time) at 888 First Street, NE., Room 2A, Washington, DC 20426.

From FERC's Home Page on the Internet, this information is available in both the Federal Energy Regulatory Records Information System (FERRIS) and the Records and Information Management System (RIMS).

—FERRIS provides access to the texts of formal documents issued by the Commission since November 14, 1994.

—FERRIS can be accessed using the FERRIS link or the Energy Information Online icon. The full text of this document is available on FERRIS in ASCII and WordPerfect 8.0 format for viewing, printing, and/or downloading.

—RIMS contains images of documents submitted to and issued by the Commission after November 16, 1981. Documents from November 1995 to the present can be viewed and printed from FERC's Home Page using the RIMS link or the Energy Information Online icon. Descriptions of documents back to November 16, 1981, are also available from RIMS-on-the-Web; requests for copies of these and other older documents should be submitted to the Public Reference Room.

User assistance is available for RIMS, FERRIS, and the Website during normal business hours from our Help line at (202) 208–2222 (E-Mail to *WebMaster@ferc.gov*) or the Public Reference at (202) 208–1371 (E-Mail to *public.referenceroom@ferc.gov*).

During normal business hours, documents can also be viewed and/or printed in FERC's Public Reference Room, where RIMS, FERRIS, and the FERC Website are available. User assistance is also available.

VIII. Effective Date and Congressional Notification

This Final Rule will take effect immediately upon issuance. Pursuant to 5 U.S.C. 804(3)(A), agencies are not required to notify Congress of any Final Rule that is a rule of particular applicability, including a rule that approves or prescribes rates, services, corporate or financial structures, reorganizations, or accounting practices. The Commission finds that this Final Rule is covered by the exception. The only impact of the rule is to prescribe the rate that the Commission's Public Reference Room contractor can charge for hard copies of certain documents. It is therefore a rule of particular applicability prescribing a rate, and the provisions of 5 U.S.C. 801 regarding Congressional review of Final Rules do not apply.

The Commission is issuing this as a final rule without a period for public comment. Under 5 U.S.C. 553(b), notice and comment procedures are unnecessary where a rulemaking concerns only agency procedure and practice, or where the agency finds that notice and comment is unnecessary. This rule concerns only matters of agency procedure and will not significantly affect regulated entities or the general public. Therefore, the Commission finds notice and comment procedures to be unnecessary.

In addition, in accordance with 5 U.S.C. 553(d)(3), the Commission finds that good cause exists to make this Final Rule effective immediately upon issuance. The increase in copying charges is necessary to make it economically viable for the Commission's Public Reference Room to continue offering this service, and will have minimal impact upon the public.

List of Subjects in 18 CFR Part 388

Confidential business information, Freedom of information.

By the Commission.

Linwood A. Watson, Jr.,

Deputy Secretary.

In consideration of the foregoing, the Commission amends Part 388, Chapter I, Title 18, of the *Code of Federal Regulations* as follows:

[4] U.S.C. 601–612.

[5] Order No. 486, 52 FR 47897 (Dec. 17, 1987); FERC Stats. & Regs. [Regulations Preambles 1986–1990] ¶ 30,783 (Dec. 10, 1984) (*codified* at 18 CFR part 380).

[6] 18 CFR 380.4(1) and (5).

[7] 5 CFR part 1320.

PART 388—INFORMATION AND REQUESTS

1. The authority citation for part 388 continues to read as follows:

Authority: 5 U.S.C. 301–305, 551, 552 (as amended), 553–557; 42 U.S.C. 7101–7352.

2. In § 388.109, paragraph (a)(4)(i) is revised to read as follows:

* * * * *

§ 388.109 Fees for record requests.

(a) * * *

(4)(i) The public may purchase hard copies of documents available in electronic form from the Commission's Federal Energy Regulatory Records Information System (FERRIS) for 20 cents per page.

[FR Doc. 02–10808 Filed 5–1–02; 8:45 am]

BILLING CODE 6717–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 520

Oral Dosage Form New Animal Drugs; Change of Sponsor

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect a change of sponsor for three approved abbreviated new animal drug applications (ANADAs) from Blue Ridge Pharmaceuticals, Inc., to Virbac AH, Inc.

DATES: This rule is effective May 2, 2002.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV–102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0209, e-mail: lluther@cvm.fda.gov.

SUPPLEMENTARY INFORMATION: Blue Ridge Pharmaceuticals, Inc., 4249–105 Piedmont Pkwy., Greensboro, NC 27410, has informed FDA that it has transferred ownership of, and all rights and interest in, NADA 200–270 for IVERHART (ivermectin) Tablets, NADA 200–281 for WORMEXX (pyrantel pamoate) Chewable Tablets, and NADA 200–302 for IVERHART Plus (ivermectin/pyrantel pamoate) Flavored Chewable Tablets to Virbac AH, Inc., 3200 Meacham Blvd., Ft. Worth, TX 76137. Accordingly, the agency is amending the regulations in 21 CFR 520.1193, 520.1196, and 520.2041 to reflect the transfer of ownership.

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.

§ 520.1193 [Amended]

2. Section 520.1193 *Ivermectin tablets and chewables* is amended in paragraph (b)(2) by removing ''065274'' and by adding in its place ''051311''.

§ 520.1196 [Amended]

3. Section 520.1196 *Ivermectin and pyrantel pamoate chewable tablet* is amended in the section heading by removing ''tablet'' and by adding in its place ''tablets''; and in paragraph (b) by removing ''065274'' and by adding in its place ''051311''.

§ 520.2041 [Amended]

4. Section 520.2041 *Pyrantel pamoate chewable tablets* is amended in paragraph (b) by removing ''065274'' and by adding in its place ''051311''.

Dated: April 3, 2002..

Andrew J. Beaulieu,

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

[FR Doc. 02–10793 Filed 5–1–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

New Animal Drugs for Use in Animal Feeds; Tilmicosin

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 Elanco Animal Health. The supplemental NADA provides for additions to labeling of tilmicosin for use in swine feed.

DATES: This rule is effective May 2, 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.

SUPPLEMENTARY INFORMATION: Elanco Animal Health, A Division of Eli Lilly & Co., Lilly Corporate Center, Indianapolis, IN 46285, filed a supplement to NADA 141–064 that provides for the use of PULMOTIL (tilmicosin phosphate) Type A medicated article in swine feed for the control of swine respiratory disease associated with certain bacterial organisms. The supplemental NADA provides for additional use information in labeling. The supplemental NADA is approved as of November 15, 2001, and the regulations are amended in 21 CFR 558.618 to reflect the approval.

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 this 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.

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 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: