Amount in controversy	
$0–9,999. (18 CFR 381.303(b))	100
10,000–29,999. (18 CFR 381.303(b))	600
30,000 or more. (18 CFR 381.303(a))	24,140
3. Review of a Department of Energy denial of adjustment:	
Amount in controversy	
$0–9,999. (18 CFR 381.304(b))	100
10,000–29,999. (18 CFR 381.304(b))	600
30,000 or more. (18 CFR 381.304(a))	12,650
4. Written legal interpretations by the Office of General Counsel. (18 CFR 381.305(a))	4,740
Fees Applicable to Natural Gas Pipelines	
1. Pipeline certificate applications pursuant to 18 CFR 284.224. (18 CFR 381.207(b))	[1] 1,000
Fees Applicable to Cogenerators and Small Power Producers	
1. Certification of qualifying status as a small power production facility. (18 CFR 381.505(a))	14,220
2. Certification of qualifying status as a cogeneration facility. (18 CFR 381.505(a))	16,090
3. Applications for exempt wholesale generator status. (18 CFR 381.801)	970

[1] This fee has not been changed.

List of Subjects in 18 CFR Part 381

Electric power plants, Electric utilities, Natural gas, Reporting and recordkeeping requirements.

Thomas R. Herlihy,

Executive Director and Chief Financial Officer.

In consideration of the foregoing, the Commission amends Part 381, Chapter I, Title 18, Code of Federal Regulations, as set forth below.

PART 381—FEES

1. The authority citation for Part 381 continues to read as follows:

Authority: 15 U.S.C. 717–717w; 16 U.S.C. 791–828c, 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.

§ 381.302 [Amended]

2. In § 381.302, paragraph (a) is amended by removing ''$ 15,760'' and adding ''$ 16,530'' in its place.

§ 381.303 [Amended]

3. In 381.303, paragraph (a) is amended by removing ''$ 23,010'' and adding ''$ 24,140'' in its place.

§ 381.304 [Amended]

4. In § 381.304, paragraph (a) is amended by removing ''$ 12,060'' and adding ''$ 12,650'' in its place.

§ 381.305 [Amended]

5. In § 381.305, paragraph (a) is amended by removing ''$ 4,520'' and adding ''$ 4,740'' in its place.

§ 381.403 [Amended]

6. Section 381.403 is amended by removing ''$ 7,840'' and adding ''$ 8,230'' in its place.

§ 381.505 [Amended]

7. In § 381.505, paragraph (a) is amended by removing ''$ 13,550'' and adding ''$ 14,220'' in its place and by removing ''$ 15,340'' and adding ''$ 16,090'' in its place.

§ 381.801 [Amended]

8. Section 381.801 is amended by removing ''$ 1,310'' and adding ''$ 970'' in its place.

[FR Doc. 01–30125 Filed 12–4–01; 8:45 am]

BILLING CODE 6717–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 510

New Animal Drugs; Change of Sponsor's Address

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's address for Merial Ltd.

DATES: This rule is effective December 5, 2001.

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.

SUPPLEMENTARY INFORMATION: Merial Ltd., 2100 Ronson Rd., Iselin, NJ 08830–3077, has informed FDA of a change of sponsor's address to 3239 Satellite Blvd., Bldg. 500, Duluth, GA 30096–4640. Accordingly, the agency is amending the regulations in 21 CFR 510.600(c)(1) and (c)(2) to reflect the change.

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 510

Administrative practice and procedure, Animal drugs, Labeling, Reporting and recordkeeping requirements.

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 510 is amended as follows:

PART 510—NEW ANIMAL DRUGS

1. The authority citation for 21 CFR part 510 continues to read as follows:

Authority: 21 U.S.C. 321, 331, 351, 352, 353, 360b, 371, 379e.

2. Section 510.600 is amended in the table in paragraph (c)(1) by revising the entry for ''Merial Ltd.'' and in the table in paragraph (c)(2) by revising the entry for ''050604'' to read as follows:

§ 510.600 Names, addresses, and drug labeler codes of sponsors of approved applications.

* * * * *

(c) * * *

(1) * * *

Firm name and address	Drug labeler code
* * * * * * *	
Merial Ltd., 3239 Satellite Blvd., Bldg. 500, Duluth, GA 30096–4640..	050604
* * * * * * *	

(2) * * *

Drug labeler code	Firm name and address
* * * * * * *	
050604	Merial Ltd., 3239 Satellite Blvd., Bldg. 500, Duluth, GA 30096–4640.
* * * * * * *	

Dated: November 15, 2001.

Claire M. Lathers,

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

[FR Doc. 01–30038 Filed 12–4–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510 and 524

Ophthalmic and Topical Dosage Form New Animal Drugs; Ivermectin Pour-On

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 Virbac AH, Inc. 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 December 5, 2001.

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: Virbac AH, Inc., 3200 Meacham Blvd., Ft. Worth, TX 76137, filed ANADA 200–318 for VIRBAMEC (ivermectin) Pour-On. The ANADA 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. Virbac's VIRBAMEC Pour-On is approved as a generic copy of Merial Ltd.'s IVOMEC Pour-On for Cattle, approved under NADA 140–841. The ANADA 200–318 is approved as of September 21, 2001, and the regulations in 21 CFR 524.1193 are amended to reflect the approval. The basis of approval is discussed in the freedom of information summary.

In addition, Virbac AH, Inc., has not been previously listed in the animal drug regulations as a sponsor of an approved application. At this time, 21 CFR 510.600(c) is being amended to add entries for the firm.

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

21 CFR Part 510

Administrative practice and procedure, Animal drugs, Labeling, Reporting and recordkeeping requirements.

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 parts 510 and 524 are amended as follows:

PART 510—NEW ANIMAL DRUGS

1. The authority citation for 21 CFR part 510 continues to read as follows:

Authority: 21 U.S.C. 321, 331, 351, 352, 353, 360b, 371, 379e.

2. Section 510.600 is amended in the table in paragraph (c)(1) by alphabetically adding an entry for ''Virbac AH, Inc.'' and in the table in paragraph (c)(2) by numerically adding an entry for ''051311'' to read as follows:

§ 510.600 Names, addresses, and drug labeler codes of sponsors of approved applications.

* * * * *

(c) * * *

(1) * * *