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Ali Bahrami,

Manager, Transport Airplane Directorate, Aircraft Certification Service.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 524

Ophthalmic and Topical Dosage Form New Animal Drugs; Gentamicin Sulfate, Betamethasone Valerate, Clotrimazole Ointment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

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 Med-Pharmex, Inc. The supplemental ANADA provides for a new container size, a 15-gram bottle, from which gentamicin sulfate, betamethasone valerate, clotrimazole ointment may be dispensed for the treatment of acute and chronic canine otitis externa.

DATES: This rule is effective April 3, 2006.

FOR FURTHER INFORMATION CONTACT: Christopher Melluso, Center for Veterinary Medicine (HFV–104), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0169, e-mail: *christopher.melluso@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: Med-Pharmex, Inc., 2727 Thompson Creek Rd., Pomona, CA 91767–1861, filed a supplement to ANADA 200–229 that provides for use of TRI-OTIC (gentamicin sulfate, USP; betamethasone valerate, USP; and clotrimazole, USP) Ointment for the treatment of canine otitis externa associated with yeast (*Malassezia pachydermatis*, formerly *Pityrosporum canis*) and/or bacteria susceptible to gentamicin. The supplement provides for a new container size, a 15-gram bottle. The supplemental ANADA is approved as of February 27, 2006, and the regulations are amended in § 524.1044g (21 CFR 524.1044g) to reflect the approval. The basis of approval is discussed in the freedom of information summary.

In addition, FDA has noticed that a 215-gram bottle size was approved for this product under ANADA 200–229 but not codified. At this time, that bottle size is being added to § 524.1044g. This action is being taken to improve the accuracy of the regulations.

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.

■ 2. In § 524.1044g, revise paragraphs (b)(2) and (c)(1)(i) to read as follows:

§ 524.1044g Gentamicin sulfate, betamethasone valerate, clotrimazole ointment.

* * * * *

(b) * * *

(2) No. 054925 for use of 7.5- or 15-g tubes; 10-, 15-, 25-, or 215-g bottles.

* * * * *

(c) * * *

(1) * * *

(i) From 7.5- or 15-g tubes; 10-, 12.5-, 15-, 25-, or 30-g bottles: 4 drops for dogs weighing less than 30 pounds (lb) or 8 drops for dogs weighing 30 lb or more.

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Dated: March 24, 2006.

Bernadette A. Dunham,

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

[FR Doc. 06–3149 Filed 3–31–06; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF STATE

22 CFR Part 34

[Public Notice: 5310]

Debt Collection

AGENCY: Department of State.

ACTION: Final rule.

SUMMARY: The Department of State is revising its debt collection regulations to conform to changes in Federal Government-wide debt collection laws and regulations, including the Debt Collection Improvement Act of 1996 and the revised Federal Claims Collections Standards jointly issued by the Department of the Treasury and Department of Justice in 2000.

DATES: This rule is effective May 3, 2006.

ADDRESSES: You may submit comments by any of the following methods:

- E-mail: *AmoryER@state.gov*
- Mail paper submissions to the Deputy Assistant Secretary for Global Financial Services, Charleston Financial Service Center, P.O. Box 150008, Charleston, S.C. 29415–5008.

Persons with access to the internet may also view this notice by going to the regulations.gov Web site at: *http://www.regulations.gov/index.cfm.*

FOR FURTHER INFORMATION CONTACT: Elizabeth Amory, Office of the Legal Adviser, Charleston Financial Service Center, telephone 843–308–5605.

SUPPLEMENTARY INFORMATION: This rule revises the Department of State's (State) debt collection regulations found at 22 CFR part 34 to conform to the Debt Collection Improvement Act of 1996 (DCIA), Public Law 104–134, 110 Stat. 1321, 1358 (Apr. 26, 1996), the revised Federal Claims Collection Standards, 31 CFR Chapter IX (parts 900 through 904), and other laws applicable to the collection of non-tax debt owed to the Government.

This regulation provides procedures for the collection of debts owed to State entities. State adopts the Government-wide debt collection standards promulgated by the Departments of the Treasury and Justice, known as the Federal Claims Collection Standards (FCCS), as revised on November 22, 2000 (65 FR 70390), and supplements the FCCS by prescribing procedures