

By the Commission.

Dated: June 21, 2002.

Margaret H. McFarland,

Deputy Secretary.

[FR Doc. 02-16211 Filed 6-26-02; 8:45 am]

BILLING CODE 8010-01-P

DEPARTMENT OF THE TREASURY

Customs Service

19 CFR PART 12

[T.D. 02-30]

RIN 1515-AD12

Extension of Import Restrictions Imposed on Archaeological and Ethnological Materials From Peru; Correction

AGENCY: Customs Service, Treasury.

ACTION: Final rule; correction.

SUMMARY: This document contains corrections to the final rule (T.D. 02-30) that was published in the **Federal Register** on June 6, 2002. The final rule extended for a period of five years from June 9, 2002, the import restrictions that were already in place for certain archaeological and ethnological materials from Peru. This document corrects the Internet web site address for accessing the Designated List of Archaeological and Ethnological Materials from Peru to which the import restrictions apply and an accompanying image database. The document also clarifies that the beginning date of the five year extension is June 9, 2002.

EFFECTIVE DATE: June 9, 2002.

FOR FURTHER INFORMATION CONTACT: (Regulatory Aspects) Joseph Howard, Intellectual Property Rights Branch (202) 927-2336; (Operational Aspects) Al Morawski, Trade Operations (202) 927-0402.

SUPPLEMENTARY INFORMATION:

Background

A final rule document, published as T.D. 02-30 in the **Federal Register** (67 FR 38877) on Thursday June 6, 2002, extended for a period of five years from June 9, 2002, the import restrictions that were already in place for certain archaeological and ethnological materials from Peru. The final rule amended section 12.104g(a), Customs Regulations (19 CFR 12.104g(a)).

This document corrects an error in the Background section of the document regarding the Internet web site address that was set forth to enable the public to access the Designated List of Archaeological and Ethnological

Materials from Peru, which describes the materials covered by the import restrictions, and an accompanying image database. The document also clarifies that the beginning date of the five year extension is June 9, 2002, by changing the effective date of the regulation to June 9, 2002.

Corrections

In rule FR Doc. 02-14219, published on June 6, 2002 (67 FR 38877), make the following corrections:

1. On page 38877, in the first column, the **EFFECTIVE DATE** section should read as follows:

EFFECTIVE DATE: June 9, 2002.

2. On page 38877, in the third column, the first full sentence should read as follows:

The list and accompanying image database may also be found at the following Internet web site address: <http://exchanges.state.gov/culprop>.

Dated: June 24, 2002.

Sandra L. Bell,

Acting Assistant Commissioner, Office of Regulations and Rulings.

[FR Doc. 02-16235 Filed 6-26-02; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510 and 520

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 two approved new animal drug applications (NADAs) and an approved abbreviated new animal drug application (ANADA) from Lambert-Kay, A Division of Carter-Wallace, Inc., to Church & Dwight Co., Inc. The drug labeler code for Church & Dwight Co., Inc., is also being listed. **DATES:** This rule is effective June 27, 2002.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV-101), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301-827-0209, e-mail: lluther@cvm.fda.gov.

SUPPLEMENTARY INFORMATION: Lambert-Kay, A Division of Carter-Wallace, Inc., P.O. Box 1001, Half Acre Rd., Cranbury, NJ 08512-0181, has informed FDA that

it has transferred ownership of, and all rights and interest in, NADA 101-497 for TINY TIGER (dichlorophene/toluene) Worming Capsules, NADA 101-498 for LK (dichlorophene/toluene) Worming Capsules, and ANADA 200-028 for EVICT (pyrantel pamoate) Liquid Wormer to Church & Dwight Co., Inc., 469 North Harrison St., Princeton, NJ 08543-5297. Accordingly, the agency is amending the regulations in §§ 520.580 and 520.2043 (21 CFR 520.580 and 520.2043) to reflect the transfer of ownership.

Church & Dwight Co., Inc., has not been previously listed in the animal drug regulations as a sponsor of an approved application. Following these changes of sponsorship, Lambert-Kay is no longer the sponsor of any approved applications. Accordingly, 21 CFR 510.600(c)(1) and (c)(2) is being amended to add entries for Church & Dwight Co., Inc., and to remove the entries for Lambert-Kay. Also, § 520.2043 is being revised to reflect a current format.

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 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 parts 510 and 520 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 removing the entry for "Lambert-Kay, A Division of Carter-Wallace, Inc." and by alphabetically adding an entry for "Church & Dwight Co., Inc." and in the table in paragraph (c)(2) by removing the entry "011615" and by numerically adding an entry for "010237" 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
* * * * *	* *
Church & Dwight Co., Inc., 469 North Harrison St., Princeton, NJ 08543-5297	010237
* * * * *	* *

(2) * * *

Drug labeler code	Firm name and address
* *	* *
010237	Church & Dwight Co., Inc., 469 North Harrison St., Princeton, NJ 08543-5297
* *	* *

**PART 520—ORAL DOSAGE FORM
NEW ANIMAL DRUGS**

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

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

§ 520.580 [Amended]

4. Section 520.580 *Dichlorophene and toluene capsules* is amended by removing footnote 1 every place it appears in the section and in paragraph (b)(1) by removing “011615” and by adding in its place “010237”.

5. Section 520.2043 is revised to read as follows:

§ 520.2043 Pyrantel pamoate suspension.

(a) *Specifications.* (1) Each milliliter (mL) contains pyrantel pamoate equivalent to 50 milligrams (mg) pyrantel base.

(2) Each mL contains pyrantel pamoate equivalent to 2.27 or 4.54 mg pyrantel base.

(3) Each mL contains pyrantel pamoate equivalent to 4.54 mg pyrantel base.

(b) *Sponsors.* See sponsors in § 510.600(c) of this chapter for uses as in paragraph (d) of this section.

(1) Nos. 000069 and 059130 for use of the product described in paragraph (a)(1) as in paragraph (d)(1) of this section.

(2) Nos. 000069, 010237, and 059130 for use of the products described in paragraph (a)(2) as in paragraph (d)(2) of this section.

(3) No. 023851 for use of the product described in paragraph (a)(3) as in paragraph (d)(2) of this section.

(c) *Special considerations.* See § 500.25 of this chapter.

(d) *Conditions of use—(1) Horses and ponies.* It is used as follows:

(i) *Amount.* 3 mg per pound (lb) body weight as a single dose mixed with the

usual grain ration, or by stomach tube or dose syringe.

(ii) *Indications for use.* For the removal and control of infections from the following mature parasites: Large strongyles (*Strongylus vulgaris*, *S. edentatus*, *S. equinus*), small strongyles, pinworms (*Oxyuris*), and large roundworms (*Parascaris*).

(iii) *Limitations.* Not for use in horses and ponies to be slaughtered for food purposes. When the drug is for administration by stomach tube, it shall be labeled: “Federal law restricts this drug to use by or on the order of a licensed veterinarian.”

(2) *Dogs.* It is used as follows:

(i) *Dogs and puppies—(A) Amount.* 2.27 mg/lb body weight as a single dose in the animal’s feed bowl by itself or mixed in a small quantity of food.

(B) *Indications for use.* For the removal of large roundworms (*Toxocara canis* and *Toxascaris leonina*) and hookworms (*Ancylostoma caninum* and *Uncinaria stenocephala*).

(C) *Limitations.* Additional treatment may be required and should be confirmed by fecal examination within 2 to 4 weeks.

(ii) *Dogs, puppies, and lactating bitches after whelping—(A) Amount.* 2.27 mg/lb body weight.

(B) *Indications for use.* To prevent reinfections of *T. canis*.

(C) *Limitations.* Administer to puppies at 2, 3, 4, 6, 8, and 10 weeks of age. Administer to lactating bitches 2 to 3 weeks after whelping. Adult dogs kept in heavily contaminated quarters may be treated at monthly intervals.

Dated: May 24, 2002.

Andrew J. Beaulieu,

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

[FR Doc. 02-16050 Filed 6-26-02; 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; 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 supplemental new animal drug application (NADA) filed by Alpharma, Inc. The supplemental NADA provides for the administration of Type C medicated feeds containing chlortetracycline to cattle as a top dress on feed for the treatment of enteritis and pneumonia.

DATES: This rule is effective June 27, 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: jmessenh@cvm.fda.gov.

SUPPLEMENTARY INFORMATION: Alpharma, Inc., One Executive Dr., P.O. Box 1399, Fort Lee, NJ 07024, filed a supplement to NADA 48-761 for AUREOMYCIN 50, 90, or 100 (chlortetracycline) Type A medicated articles. The supplemental NADA provides for the administration of Type C medicated feeds containing chlortetracycline to calves, beef and nonlactating dairy cattle as a top dress on feed to deliver 10 milligrams (mg) chlortetracycline per pound of body weight daily. These medicated feeds are used for the treatment of bacterial enteritis caused by *Escherichia coli* and