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DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

7 CFR Parts 318 and 319

[Docket No. 02–019–2]

Phytosanitary Treatments; Location of Treatment Schedules and Other Requirements; Correction

AGENCY: Animal and Plant Health Inspection Service, USDA.

ACTION: Final rule; correction.

SUMMARY: We are correcting errors in the amendatory instructions in our final rule that removed the Plant Protection and Quarantine Treatment Manual from the list of materials incorporated by reference and added treatment schedules and related requirements from that document to our phytosanitary treatments regulations. The final rule was effective and published in the **Federal Register** on June 7, 2005 (70 FR 33263–33326, Docket No. 02–019–1).

DATES: Effective July 15, 2005.

FOR FURTHER INFORMATION CONTACT: Ms. Meredith C. Jones, Regulatory Coordination Specialist, PPQ, APHIS, 4700 River Road Unit 141, Riverdale, MD 20737–1236; (301) 734–7467.

SUPPLEMENTARY INFORMATION: In a final rule effective and published in the **Federal Register** on June 7, 2005 (70 FR 33263–33326, Docket No. 02–019–1), we amended the plant health regulations by adding to 7 CFR part 305 treatment schedules and related requirements that had appeared in the Plant Protection and Quarantine Treatment Manual and by removing the Plant Protection and Quarantine Treatment Manual from the list of materials is incorporated by reference into the regulations.

In the final rule, it was our intention to amend the regulations by, among other things:

• In § 318.58–4a, redesignating paragraphs (b) through (d) as paragraphs (a) through (c), respectively;

• Revising the newly redesignated paragraph § 318.58–4a(a);

• In § 319.37–6(d)(2), redesignating footnote 9 as footnote 8 and revising the newly redesignated footnote 8;

• Amending § 319.56–2h; and

• In § 319.56–2h(d), removing the words ''the Plant Protection and Quarantine Treatment Manual'' and adding the words ''part 305 of this chapter'' in their place.

However, our amendatory instructions that were intended to accomplish these changes were either absent or erroneous. This document corrects these errors.

■ In FR Doc. 05–9387, published on June 7, 2004 (70 FR 33263–33326), make the following corrections:

§ 318.58–4a [Corrected]

■ 1. On page 33324, in the amendments to 7 CFR part 318, in instruction 34 for § 318.58–4a, instruction b. is corrected to read as follows, and a new instruction d. is added to read as follows:

■ b. By redesignating paragraphs (b) through (d) as paragraphs (a) through (c), respectively.

■ d. By revising newly redesignated paragraph (a) to read as set forth below.

§ 319.37–6 [Corrected]

■ 2. On page 33324, in the third column, in § 319.37–6, footnote number 3 is corrected to read footnote number 8.

§ 319.56–2h [Corrected]

■ 3. On page 33325, in the amendments to 7 CFR part 319, instruction 58 for § 319.56–2h is corrected to read as follows, and in instruction 58, instruction c. is corrected to read as follows:

■ 58. Section 319.56–2h is amended as follows:

■ c. In paragraph (d), by removing the words ''the Plant Protection and Quarantine Treatment Manual'' and adding the words ''part 305 of this chapter'' in their place.

Done in Washington, DC, this 7th day of July, 2005.

Elizabeth E. Gaston,

Acting Administrator, Animal and Plant Health Inspection Service.

[FR Doc. 05–13853 Filed 7–14–05; 8:45 am]

BILLING CODE 3410–34–P

NUCLEAR REGULATORY COMMISSION

10 CFR Part 72

RIN 3150–AH72

List of Approved Spent Fuel Storage Casks: Standardized NUHOMS®–24P, –52B, –61BT, –32PT, –24PHB, and –24PTH Revision; Withdrawal of Direct Final Rule

AGENCY: Nuclear Regulatory Commission.

ACTION: Direct final rule; withdrawal.

SUMMARY: The Nuclear Regulatory Commission (NRC) is withdrawing a direct final rule that would have revised the NUHOMS®–24P, –52B, –61BT, –32PT, –24PHB, and –24PTH cask system listing within the list of approved spent fuel storage casks to include Amendment No. 8 to the Certificate of Compliance (CoC). The NRC is taking this action because the NRC staff has become aware of changes in the Technical Specifications (TS) associated with this CoC. A notice withdrawing the companion proposed rule is published in the proposed rule section of this **Federal Register**.

FOR FURTHER INFORMATION CONTACT: Jayne M. McCausland, Office of Nuclear Material Safety and Safeguards, U.S. Nuclear Regulatory Commission, Washington, DC 20555, telephone (301) 415–6219 (e-mail: *jmm2@nrc.gov*).

SUPPLEMENTARY INFORMATION: On May 25, 2005 (70 FR 29931), the NRC published in the **Federal Register** a direct final rule amending its regulations in 10 CFR 72.214 to revise the Standardized NUHOMS® System listing within the ''List of Approved Spent Fuel Storage Casks'' to include Amendment No. 8 to the CoC. Amendment No. 8 modifies the present cask system by adding a new spent fuel storage and transfer system, designated the NUHOMS®–24PTH System. The direct final rule was to become effective on August 8, 2005. The NRC also concurrently published a companion proposed rule on May 25, 2005 (70 FR 30015).

The NRC has become aware of changes in the TS associated with this CoC; therefore, the NRC is withdrawing the direct final rule. The NRC will publish a direct final rule, and its

companion proposed rule, after the needed revisions to the TS are made.

Dated at Rockville, Maryland, this 6th day of July, 2005.

For the Nuclear Regulatory Commission.

Martin J. Virgilio,

Acting Executive Director for Operations.

[FR Doc. 05–13933 Filed 7–14–05; 8:45 am]

BILLING CODE 7590–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Chapter I

[Docket No. 2005N–0201]

Change of Name and Address; Technical Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending its regulations to reflect a change in the name and address for the Association of Official Analytical Chemists International (AOAC). This action is editorial in nature and is intended to improve the accuracy of the agency's regulations.

DATES: This rule is effective July 15, 2005.

FOR FURTHER INFORMATION CONTACT: Joyce Strong, Office of Policy and Planning (HF–27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7010.

SUPPLEMENTARY INFORMATION: This document amends FDA's regulations to reflect the name and address change of AOAC by removing the outdated name and address wherever it appears and by adding the new name and address in its place in 21 CFR parts 2, 10, 101, 102, 106, 114, 130, 131, 133, 135, 136, 137, 139, 145, 146, 150, 155, 156, 160, 161, 163, 164, 166, 168, 169, 172, 173, 176, 177, 178, 184, 189, 211, 226, 520, and 573.

Publication of this document constitutes final action on these changes under the Administrative Procedure Act (5 U.S.C. 553). Notice and public procedure are unnecessary because FDA is merely correcting nonsubstantive errors.

■ Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR chapter I is amended as follows:

Chapter I [Nomenclature changes]

■ 1. Parts 2, 101, 102, 106, 114, 130, 131, 133, 135, 136, 137, 139, 145, 146, 150, 155, 156, 160, 161, 163, 164, 166, 168, 169, 172, 173, 176, 177, 178, 184, 189, 211, 226, 520, and 573 are amended by removing the text set forth below wherever it appears and adding new text in its place as follows:

■ A. Remove:

''Association of Official Analytical Chemists International, 481 North Frederick Ave., suite 500, Gaithersburg, MD 20877–2504'', or

''Association of Official Analytical Chemists International, 481 North Frederic Ave., suite 500, Gaithersburg, MD 20877–2504'', or

''Association of Official Analytical Chemists International, 481 North Frederick Ave., suite 500, Gaithersbuerg, MD 20877–2504'', or

''Association of Official Analytical Chemists International, 481 North Frederick Ave., Suite 500, Gaithersburg, MD 20877–2504'', or

''Association of Official Analytical Chemists International, 481 North Frederick Ave, suite 500, Gaithersburg, MD 20877–2504'', or

''AOAC INTERNATIONAL, 481 North Frederick Ave., suite 500, Gaithersburg, MD 20877–2504'', or

''Association of Official Analytical Chemists, 2200 Wilson Blvd., Suite 400, Arlington, VA 22201–3301'', or

''Association of Official Analytical Chemists, P.O. Box 540, Benjamin Franklin Station, Washington, DC 20044''.

■ B. Add:

''Association of Official Analytical Chemists International, 481 North Frederick Ave., suite 500, Gaithersburg, MD 20877''.

PARTS 2, 10, 101, and 211 [AMENDED]

■ 2. In addition to the amendments set forth in the previous paragraph, in 21 CFR parts 2, 10, 101, and 211 add the word ''International'' after the words ''Association of Official Analytical Chemists'' in the following places:

a. Section 2.19 where it appears in the first sentence, after the words ''to utilize the methods of analysis of the'';

b. Section 10.95(d)(8)(v);

c. Section 101.70(f);

d. Section 101.81(c)(2)(ii)(B)(*2*) in the first sentence;

e. Appendix A to part 101; and

f. Section 211.194(a)(2) in the third sentence.

Dated: July 8, 2005.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 05–13898 Filed 7–14–05; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 520

Oral Dosage Form New Animal Drugs; Lincomycin and Spectinomycin Soluble Powder

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 Cross Vetpharm Group Ltd. The ANADA provides for the oral use of lincomycin and spectinomycin soluble powder to create a solution administered in the drinking water of chickens as an aid in the control of airsacculitis.

DATES: This rule is effective July 15, 2005.

FOR FURTHER INFORMATION CONTACT: Daniel A. Benz, Center for Veterinary Medicine (HFV–104), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0223, e-mail: *daniel.benz@fda.gov*.

SUPPLEMENTARY INFORMATION: Cross Vetpharm Group Ltd., Broomhill Rd., Tallaght, Dublin 24, Ireland, filed ANADA 200–380 that provides for use of SPECLINX–50 (spectinomycin dihydrochloride pentahydrate and lincomycin hydrochloride monohydrate) Water Soluble Powder to create a solution administered in the drinking water of chickens. This solution acts as an aid in the control of airsacculitis caused by either *Mycoplasma synoviae* or *M. gallisepticum* susceptible to lincomycin-spectinomycin and complicated chronic respiratory disease (air sac infection) caused by *Escherichia coli* and *M. gallisepticum* susceptible to lincomycin-spectinomycin. Cross Vetpharm Group Ltd.'s SPECLINX–50, Water Soluble Powder is approved as a generic copy of Pharmacia & Upjohn Co.'s L-S 50 Water Soluble Powder, approved under NADA 046–109. The ANADA is approved as of June 7, 2005, and the regulations are amended in 21 CFR 520.1265 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