

technical regulations that require frequent and routine amendments to keep them operationally current. Therefore, this regulation—(1) is not a “significant regulatory action” under Executive Order 12866; (2) is not a “significant rule” under DOT Regulatory Policies and Procedures (44 FR 11034; February 26, 1979); and (3) if promulgated, will not have a significant economic impact, positive or negative, on a substantial number of small entities under the criteria of the Regulatory Flexibility Act.

List of Subjects 14 CFR Part 71

Airspace, Incorporation by reference, Navigation (air).

Adoption of the Amendment

■ Accordingly, the Federal Aviation Administration amends 14 CFR part 71 as follows:

PART 71—DESIGNATION OF CLASS A, CLASS B, CLASS C, CLASS D, AND CLASS E AIRSPACE AREAS; AIRWAYS; ROUTES; AND REPORTING POINTS

■ 1. The authority citation for 14 CFR part 71 continues to read as follows:

Authority: 49 U.S.C. 106(g), 40103, 40113, 40120; E.O. 10854; 24 FR 9565, 3 CFR, 1959–1963 Comp., p. 389.

§ 71.1 [Amended]

■ 2. The incorporation by reference in 14 CFR 71.1 of the Federal Aviation Administration Order 7400.9K, dated August 30, 2002, and effective September 16, 2002, is amended as follows:

Paragraph 6005 Class E airspace areas extending upward from 700 feet or more above the surface of the earth.

* * * * *

AWP AZ E5 Window Rock, AZ [Revised]

Window Rock Airport, AZ
(Lat. 35°39'07" N, long. 109°04'02" W)
Gallup VORTAC
(Lat. 35°28'34" N, long. 108°52'21" W)

That airspace extending upward from 700 feet above the surface within 6.6-mile radius of the Window Rock Airport and within 2.6 miles each side of the Gallup VORTAC 318° radial, extending from the 6.6-mile radius to the Gallup VORTAC and within 4-miles west and 2 miles east of the 214° bearing from the Window Rock airport, extending from the 6.6-mile radius to 13.4 miles southwest of the airport and within 2 miles each side of 004° bearing from the Window Rock Airport, extending from the 6.6-mile radius to 11.7 miles north of the airport and within an area bounded by a line beginning at Lat. 35°38'27" N, Long. 109°06'37" W; to Lat. 35°31'07" N, Long. 108°58'34" W; to Lat. 35°27'13" N, Long. 109°04'36" W; to Lat. 35°25'26" N, Long. 109°14'07" W; to lat 35°31'35" N, Long.

109°11'00" W, to the point of beginning. That airspace extending upward from 1,200 feet above the surface bounded by a line beginning at Lat. 35°30'00" N, Long. 109°17'00" W; to Lat. 35°28'00" N, Long. 109°30'00" W; to Lat. 35°08'00" N, Long. 109°39'00" W; to Lat. 35°08'00" N, Long. 109°25'00" W; to Lat. 35°20'00" N, Long. 109°12'00" W; to the point of beginning and that airspace beginning at Lat. 35°49'30" N, Long. 109°05'00" W; to Lat. 36°04'00" N, Long. 109°27'00" W; to Lat. 36°07'00" N, Long. 109°23'00" W; to Lat. 35°54'00" N, Long. 109°03'00" W; to Lat. 35°54'00" N, Long. 108°43'00" W; to Lat. 35°51'00" N, Long. 108°44'00" W; to Lat. 35°51'30" N, Long. 108°47'00" W; to Lat. 35°44'00" N, Long. 108°51'30" W, to the point of beginning.

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Issued in Los Angeles, California, on June 5, 2003.

John Clancy,

Manager, Air Traffic Division, Western-Pacific Region.

[FR Doc. 03–15526 Filed 6–18–03; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

New Animal Drugs for Use in Animal Feeds; Lasalocid; Technical Amendment

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 new animal drug application (NADA) filed by AlphaPharma, Inc. The supplemental NADA provides for the use of a lasalocid Type A medicated article to make free-choice, loose mineral Type C medicated feeds used for increased rate of weight gain in pasture cattle (slaughter, stocker, feeder cattle, and dairy and beef replacement heifers). The regulations are also being revised to provide current references for the amounts of selenium and ethylenediamine dihydroiodide (EDDI) permitted in other free-choice cattle feeds.

DATES: This rule is effective June 19, 2003.

FOR FURTHER INFORMATION CONTACT: Eric S. Dubbin, Center for Veterinary Medicine (HFV–126), Food and Drug

Administration, 7500 Standish Pl., Rockville, MD 20855; 301–827–0232; e-mail: edubbin@cvm.fda.gov.

SUPPLEMENTARY INFORMATION: AlphaPharma, Inc., One Executive Dr., P.O. Box 1399, Fort Lee, NJ 07024, filed a supplement to NADA 96–298 that provides for use of BOVATEC 68 (lasalocid) Type A medicated article to make a free-choice high phosphorus loose mineral Type C medicated feed containing 1088 grams lasalocid per ton of feed. The free-choice medicated feed is used for increased rate of weight gain in pasture cattle (slaughter, stocker, feeder cattle, and dairy and beef replacement heifers). The NADA is approved as of April 9, 2003, and the regulations are amended in 21 CFR 558.311 to reflect the approval. The basis of approval is discussed in the freedom of information summary.

Section 558.311 is also being revised to reflect publication of an updated compliance policy guide (CPG) on permitted levels of EDDI in animal feed (CPG 7125.18, May 1, 2000).

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 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.

Under section 512(c)(2)(F)(iii) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 360b(c)(2)(F)(iii)), this approval qualifies for 3 years of marketing exclusivity beginning April 9, 2003.

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 Subject 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:

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

■ 1. The authority citation for 21 CFR part 558 continues to read as follows:

Authority: 21 U.S.C. 360b, 371.

■ 2. Section 558.311 is amended:

a. In paragraph (b)(4) by removing “(e)(2) and (e)(3)” and by adding in its place “(e)(2), (e)(3), and (e)(4)”;

b. In paragraphs (e)(2)(i) and (e)(3)(i) by revising footnote 1;

c. By redesignating paragraph (e)(4) as paragraph (e)(5); and

d. By adding new paragraph (e)(4).

The revisions and addition read as follows:

§ 558.311 Lasalocid.

* * * * *

- (e) * * *
- (2) * * *
- (i) * * *

¹ Content of this vitamin and trace mineral premixes may be varied; however, they should be comparable to those used by the firm for other free-choice feeds. Formulation modifications require FDA approval prior to marketing. Selenium must comply with § 573.920 of this chapter. Ethylenediamine dihydroiodide (EDDI) should comply with FDA Compliance Policy Guide Sec. 651.100 (CPG 7125.18).

- (3) * * *
- (i) * * *

¹ Content of vitamin and trace mineral premixes may be varied; however, they should be comparable to those used for other free-choice liquid feeds. Formulation modifications require FDA approval prior to marketing. Selenium must comply with § 573.920 of this chapter. EDDI should comply with FDA Compliance Policy Guide Sec. 651.100 (CPG 7125.18).

* * * * *

(4) It is used as a free-choice, loose mineral Type C feed as follows:

(i) *Specifications.*

| Ingredient | Percent | International feed No. |
|-------------------------------------------|---------|------------------------|
| Monocalcium Phosphate (21% P) | 57.50 | 6-01-082 |
| Salt | 17.55 | 6-04-152 |
| Distillers Dried Grains w/Solubles | 5.40 | 5-28-236 |
| Dried Cane Molasses (46% Sugars) | 5.20 | 4-04-695 |
| Potassium Chloride | 4.90 | 6-03-755 |
| Trace Mineral/Vitamin Premix ¹ | 3.35 | |

| Ingredient | Percent | International feed No. |
|-----------------------------------------------------|---------|------------------------|
| Calcium Carbonate (38% Ca) | 2.95 | 6-01-069 |
| Mineral Oil | 1.05 | 8-03-123 |
| Magnesium Oxide (58% Mg) | 1.00 | 6-02-756 |
| Iron Oxide (52% Fe) | 0.10 | 6-02-431 |
| Lasalocid Type A Medicated Article (68 g per pound) | 0.80 | |

¹ Content of vitamin and trace mineral premixes may be varied. However, they should be comparable to those used for other free-choice loose mineral feeds. Formulation modifications require FDA approval prior to marketing. Selenium must comply with § 573.920 of this chapter. EDDI should comply with FDA Compliance Policy Guides Sec. 651.100 (CPG 7125.18).

(ii) *Amount.* 1,088 grams per ton.

(iii) *Indications for use.* Pasture cattle (slaughter, stocker, feeder cattle, and dairy and beef replacement heifers): For increased rate of weight gain. Intakes of lasalocid in excess of 200 mg/head/day have not been shown to be more effective than 200 mg/head/day.

(iv) *Limitations.* Feed continuously on a free-choice basis at a rate of 60 to 300 mg lasalocid per head per day.

(v) *Sponsor.* See No. 046573 in § 510.600(c) of this chapter.

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Dated: May 29, 2003.

Steven D. Vaughn,

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

[FR Doc. 03-15541 Filed 6-18-03; 8:45 am]

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DEPARTMENT OF THE INTERIOR

Indian Arts and Crafts Board

25 CFR Part 309

RIN 1076-AE16

Protection of Products of Indian Art and Craftsmanship; Correction

AGENCY: Indian Arts and Crafts Board (IACB), Department of the Interior.

ACTION: Correction to final regulations.

SUMMARY: This document contains a correction to the final regulations for the Indian Arts and Crafts Enforcement Act of 2000 (25 CFR part 309), which were published Thursday, June 12, 2003, (68 FR 35164). The rule clarifies the regulatory definition of “Indian product,” as defined under the Indian Arts and Crafts Act of 1990.

EFFECTIVE DATES: June 13, 2003.

FOR FURTHER INFORMATION CONTACT: Meridith Z. Stanton, Director, (202) 208-3773 (not a toll free call).

SUPPLEMENTARY INFORMATION:

Background

The final regulations that are the subject of this correction clarify the regulatory definition of “Indian product,” as defined under the Indian Arts and Crafts Act of 1990 (Pub. L. 101-644, 104 Stat. 4662).

Need for Correction

As published, the final regulations contain an error which may prove to be misleading and is in need of clarification.

Correction of Publication

■ Accordingly, the publication on June 12, 2003 of the final regulations (25 CFR part 309), which were the subject of FR Doc. 03-14827, is corrected as follows:

Effective Date—[Corrected]

■ On page 35164, in the second column, the effective date of September 10, 2003 is to read July 14, 2003.

Meridith Z. Stanton,

Director, Indian Arts and Crafts Board.

[FR Doc. 03-15417 Filed 6-18-03; 8:45 am]

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DEPARTMENT OF HOMELAND SECURITY

Coast Guard

33 CFR Part 165

[COTP Los Angeles-Long Beach 01-013]

RIN 1625-AA00 (Formerly RIN 2115-AA97)

Security Zone; Port Hueneme Harbor, Ventura County, CA

AGENCY: Coast Guard, DHS.

ACTION: Temporary final rule; change in effective period.

SUMMARY: The Coast Guard is revising the effective period for a temporary security zone covering all waters within Port Hueneme Harbor in Ventura County, CA. This security zone is needed for national security reasons to protect Naval Base Ventura County and commercial port from potential subversive acts. Entry into this zone is prohibited unless specifically authorized by the Capitan of the Port Los Angeles-Long Beach, the Commanding Officer of Naval Base Ventura County, or their designated representatives.