

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-miles 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.*

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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 Alpharma, 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](mailto:edubbin@cvm.fda.gov).

**SUPPLEMENTARY INFORMATION:** Alpharma, 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: