substantial number of small entities under the criteria of the Regulatory Flexibility Act.

List of Subjects in 14 CFR Part 71

Airspace, Incorporation by reference, Navigation (air).

Adoption of the Amendment

In consideration of the foregoing, 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 Federal Aviation Administration Order 7400.9H, *Airspace Designations and Reporting Points,* dated September 1, 2000, and effective September 16, 2000, is amended as follows:

* * * * *

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

* * * * *

AAL AK E5 Indian Mountain, AK [New]

Indian Mountain LRRS, AK

(lat. 65° 59′ 34″ N., long. 153° 42′ 16″ W.)

That airspace extending upward from 700 feet above the surface within a 4 mile radius of Indian Mountain LRRS; and that adjacent airspace extending upward from 1,200 feet above the surface from lat. 66° 00′ 00″ N long. 154° 05′ 00″ W, to lat. 66° 00′ 00″ N long. 153° 00′ 00″ W, to lat. 66° 09′ 00″ N long. 153° 00′ 00″ W, to lat. 66° 09′ 00″ N long. 153° 40′ 00″ W, to lat. 66° 06′ 00″ N long. 154° 00′ 00″, thence to the point of the beginning, excluding the existing Class E airspace.

* * * * *

Issued in Anchorage, AK, on January 2, 2001.

Stephen P. Creamer,

Acting Manager, Air Traffic Division, Alaskan Region.

[FR Doc. 01–701 Filed 1–9–01; 8:45 am]

BILLING CODE 4910–13–U

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 314

[Docket No. 98N–0720]

Conforming Regulations Regarding Removal of Section 507 of the Federal Food, Drug, and Cosmetic Act; Technical Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending the regulations for applications for FDA approval to market a new drug to correct inadvertent errors. This action is necessary to ensure the accuracies and consistency of the regulation.

DATES: This rule is effective January 16, 2001.

FOR FURTHER INFORMATION CONTACT: Christine F. Rogers, Center for Drug Evaluation and Research (HFD–7), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–594–2041.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of January 5, 1999 (64 FR 396), FDA published a direct final rule that removed from the agency's regulations references to the now-repealed statutory provision of the Federal Food, Drug, and Cosmetic Act (the act) under which the agency certified antibiotic drugs (conforming regulation). Section 314.430(f) (21 CFR 314.430(f)) provides that safety and effectiveness data and information in an application may be disclosed to the public when certain events happen. Prior to the conforming regulation, § 314.430(f)(6) read: ''For applications or abbreviated applications submitted under sections 505(j) and 507 of the act, when FDA sends an approval letter to the applicant''.

The conforming regulation inadvertently changed ''section 505(j)'' to ''section 505'' and failed to remove the word ''applications'' from the introductory clause the first time it appeared. This document corrects those errors. Publication of this document constitutes final action under the Administrative Procedure Act (5 U.S.C. 553). FDA has determined that notice and public comment are unnecessary because this amendment is nonsubstantive.

List of Subjects in 21 CFR Part 314

Administrative practice and procedure, Confidential business information, Drugs, Reporting and recordkeeping requirements.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR part 314 is amended as follows:

PART 314—APPLICATIONS FOR FDA APPROVAL TO MARKET A NEW DRUG

1. The authority citation for 21 CFR part 314 continues to read as follows:

Authority: 21 U.S.C. 321, 331, 351, 352, 353, 355, 371, 374, 379e.

§ 314.430 [Amended]

2. Section 314.430 *Availability for public disclosure of data and information in an application or abbreviated application* is amended in paragraph (f)(6) by removing ''applications or'' and by removing ''505'' and adding in its place ''505(j)''.

Dated: January 4, 2001.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning, and Legislation.

[FR Doc. 01–680 Filed 1–9–01; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

New Animal Drugs for Use in Animal Feeds; Decoquinate, Monensin, and Tylosin

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 new animal drug application (NADA) filed by Alpharma, Inc. The NADA provides for use of approved, single-ingredient decoquinate, monensin, and tylosin Type A medicated articles to make three-way combination drug Type B and Type C medicated feeds used for prevention of coccidiosis, improved feed efficiency, and reduction of incidence of liver abscesses in growing-finishing cattle fed in confinement for slaughter.

DATES: This rule is effective January 10, 2001.