submit their requests and supporting documents to USDA, FSA, ORAS, Attention: Phil Brockman, PO Box 23278, Washington, DC 20026–3278. All requests and supporting documents must be received by the appropriate FSA office by August 30, 2002.

(c) Each person on the county FSA office lists may participate in the sign-up period. Eligible producers must date and sign their name on the ''County FSA Office Sign-up Sheet.'' A person whose name does not appear on the county FSA office list may participate in the sign-up period. Such person must be identified on FSA–578 during the representative period or provide documentation that demonstrates that the person was a cotton producer during the representative period. Cotton producers not listed on the FSA–578 shall submit at least one sales receipt for cotton they planted during the representative period. Cotton producers must make requests to the county FSA office where the producer's farm is located. If the producer's land is in more than one county, the producer shall make request at the county office where FSA administratively maintains and processes the producer's farm records. It is the responsibility of the person to provide the information needed by the county FSA office to determine eligibility. It is not the responsibility of the county FSA office to obtain this information. If any person whose name does not appear on the county FSA office list fails to provide at least one sales receipt for the cotton they produced during the representative period, the county FSA office shall determine that such person is ineligible to participate in the sign-up period, and shall note ineligible in the remarks section next to the person's name on the county FSA office sign-up sheet. In lieu of personally appearing at a county FSA office, eligible producers may request a sign-up form from the county FSA office where the producer's farm is located. If the producer's land is in more than one county, the producer shall make the request for the sign-up form at the county office where FSA administratively maintains and processes the producer's farm records. Such request must be accompanied by a copy of at least one sales receipt for cotton they produced during the representative period. The appropriate FSA office must receive all completed forms and supporting documentation by August 30, 2002.

7. In § 1205.28, the first sentence is revised to read as follows:

§ 1205.28 Counting.

County FSA offices and FSA, Director for Operations Review and Analysis Staff (ORAS), shall begin counting requests no later than September 3, 2002. * * *

8. Section 1205.29 is revised to read as follows:

§ 1205.29 Reporting results.

(a) Each county FSA office shall prepare and transmit to the state FSA office, by September 10, 2002, a written report of the number of eligible producers who requested the conduct of a referendum, and the number of ineligible persons who made requests.

(b) ORAS shall prepare, by September 10, 2002, a written report of the number of eligible importers who requested the conduct of a referendum, and the number of ineligible persons who made requests.

(c) Each state FSA office shall, by September 17, 2002, forward all county reports to ORAS. By September 24, 2002, ORAS shall forward its report of the total number of eligible producers and importers that requested a continuance referendum, through the sign-up period, to the Deputy Administrator, Cotton Program, AMS, Stop 0224, 1400 Independence Ave., SW, Washington, DC 20250–0224.

(d) The Chief of the Research and Promotion Staff, Cotton Program, shall prepare a report of the requests received, including the number of eligible persons who requested the conduct of a referendum, and the number of ineligible persons who made requests, to the Deputy Administrator of the Cotton Program, and shall maintain one copy of the report where it will be available for public inspection for a period of 5 years following the end of the sign-up period.

(e) The Deputy Administrator of the Cotton Program shall prepare and submit to the Secretary a report of the results of the sign-up period. The Secretary will conduct a referendum if requested by 10 percent or more of the number of cotton producers and importers voting in the most recent (July 1991) referendum, but not more than 20 percent of the total requests counted toward the 10 percent figure may be from producers in any one state or from importers of cotton. The Secretary shall announce the results of the sign-up period in a separate notice in the **Federal Register**.

Dated: April 25, 2002.

A.J. Yates,

Administrator, Agricultural Marketing Service.

[FR Doc. 02–10676 Filed 4–26–02; 11:17 am]

BILLING CODE 3410–02–P

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 71

[Airspace Docket No. 99–AWP–24]

Modification of Class E Airspace; Daggett, CA

AGENCY: Federal Aviation Administration (FAA), DOT.

ACTION: Final rule.

SUMMARY: This action modifies the Class E airspace area at Daggett, CA. The establishment of an Area navigation (RNAV) Global Positing System (GPS) Standard Instrument Approach Procedure (SIAP) RNAV (GPS) Runway (RWY) 22 SIAP and a RNAV (GPS) RWY 26 SIAP to Daggett Airport, Barstow-Daggett Airport, CA has made this action necessary. Additional controlled airspace extending upward from 700 feet or more above the surface of the earth is needed to contain aircraft executing the RNAV (GPS) RWY 22 SIAP and RNAV (GPS) 26 SIAP to Barstow-Daggett Airport. The intended effect of this action is to provide adequate controlled airspace for Instrument Flight Rules operations at Barstow-Daggett Airport, Daggett, CA.

EFFECTIVE DATE: 0901 UTC June 13, 2002.

FOR FURTHER INFORMATION CONTACT: Jeri Carson, Airspace Specialist, Airspace Branch, AWP–520, Air Traffic Division, Western-Pacific Region, Federal Aviation Administration, 1500 Aviation Boulevard, Lawndale, California 90261, telephone (310) 725—6611.

SUPPLEMENTARY INFORMATION:

History

On February 6, 2002, the FAA proposed to amend 14 CFR part 71 by modifying the Class E airspace area at Daggett, CA (67 FR 5528). Additional controlled airspace extending upward from 700 feet or more above the surface is needed to contain aircraft executing the RNAV (GPS) RWY 22 SIAP and RNAV (GPS) 26 SIAP to Barstow-Daggett Airport. This action will provide adequate controlled airspace for aircraft executing the RNAV (GPS) RWY

22 SIAP and RNAV (GPS) 26 SIAP to Barstow-Daggett Airport, Daggett, CA.

Interested parties were invited to participate in this rulemaking, proceeding by submitting written comments on the proposed to the FAA. No comments to the proposal were received. Class E airspace designations for airspace extending upward from 700 feet or more above the surface of the earth are published in paragraph 6005 of FAA Order 7400.9J, dated August 31, 2001, and effective September 16, 2001, which is incorporated by reference in 14 CFR 71.1. The Class E airspace designation listed in this document will be published subsequently in the Order.

The Rule

This amendment to 14 CFR part 71 modifies the Class E airspace area at Daggett, CA. The establishment of a RNAV (GPS) RWY 22 SIAP and RNAV RWY 26 SIAP to Barstow-Daggett Airport has made this action necessary. The effect of this action will provide adequate airspace for aircraft executing the RNAV (GPS) RWY 22 SIAP and RNAV (GPS) RWY 26 SIAP to Barstow-Daggett Airport, Daggett, CA.

The FAA has determined that this regulation only involves an established body of technical regulations for which frequent and routine amendments are necessary to keep them operationally current. Therefore, this regulation—(1) is not a ''significant regulatory action'' under Executive Order 12766; (2) is not a ''significant rule'' under DOT Regulatory Policies and Procedures (44 FR 11034; February 26, 1979); and (3) does not warrant preparation of a Regulatory Evaluation as the anticipated impact is so minimal. Since this is a routine matter that will only affect air traffic procedures and air navigation, it is certified that this rule will not have a significant economic impact on a 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; 14 CFR 11.69.]

§ 71.1 [Amended]

2. The incorporation by reference in 14 CFR 71.1 of the Federal Aviation Administration Order 7400.9J, Airspace Designations and Reporting Points, dated August 31, 2001, and effective September 16, 2001, is amended as follows:

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

* * * * *

AWP CA E5 Daggett, CA [Revised]

Barstow-Daggett Airport, CA
(Lat. 34°51′13″ N, long. 116°47′12″ W)

That airspace extending upward from 700 feet above the surface within a 6.5-mile radius of the Barstow-Daggett Airport and within 2.2 miles each side of the 057° bearing from the Barstow-Daggett Airport extending from the 6.5-mile radius to 11.8 miles northeast of the airport.

Issued in Los Angeles, California, on April 8, 2002.

John Clancy,

Manager, Air Traffic Division, Western-Pacific Region.

[FR Doc. 02–10499 Filed 4–29–02; 8:45 am]

BILLING CODE 4910–13–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

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 an approved new animal drug application (NADA) from Boehringer Ingelheim Vetmedica, Inc., to Alpharma, Inc.

DATES: This rule is effective April 30, 2002.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV–102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0209, e-mail: lluther@cvm.fda.gov.

SUPPLEMENTARY INFORMATION: Boehringer Ingelheim Vetmedica, Inc., 2621 North Belt Hwy., St. Joseph, MO 64506–2002, has informed FDA that it has transferred ownership of, and all rights and interest in, NADA 39–077 for CSP (chlortetracycline, sulfathiazole, penicillin) 250 and CSP 500 Type A medicated articles to Alpharma, Inc., One Executive Dr., P.O. Box 1399, Fort Lee, NJ 07024. Accordingly, the agency is amending the regulations in 21 CFR 558.155 to reflect the change of sponsor.

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 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.

§ 558.155 [Amended]

1. Section 558.155 *Chlortetracycline, sulfathiazole, penicillin* is amended in paragraphs (a)(1) and (a)(2) by removing ''Nos. 000010 and 046573'' and by adding in its place ''No. 046573''.

Dated: February 22, 2002.

Claire M. Lathers,

Director, Office of New Animal Drugs, Center for Veterinary Medicine.

[FR Doc. 02–10511 Filed 4–29–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 888

[Docket No. 99P–1864]

Orthopedic Devices: Reclassification of the Hip Joint Metal/Polymer Constrained Cemented or Uncemented Prosthesis

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is reclassifying the hip joint metal/polymer constrained cemented or uncemented prosthesis intended to replace a hip joint from class III (premarket approval) to class II (special controls). FDA is also