were received. Accordingly, the amendment is adopted as proposed.

In addition, the airspace docket number appearing in the Notice of Proposed Rulemaking erroneously read "88–ANM–12." This action corrects it to read "89–ANM–12."

Section 71.181 of part 71 of the Federal Aviation Regulations was republished in FAA Handbook 7400.6E, January 3, 1989.

The Rule

This amendment to part 71 of the Federal Aviation Regulations alters the McCall, ID, transition area to provide additional controlled airspace for aircraft executing a new instrument approach procedure to the McCall Municipal Airport. This alteration is intended to ensure segregation of aircraft operating under Instrument Flight Rules from aircraft operating under Visual Flight Rules. The transition area, as altered, will be depicted on aeronautical charts for pilot reference.

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. It, therefore—(1) is not a "major rule" under Executive Order 12291; (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, positive or negative, on a substantial number of small entities under the criteria of the Regulatory Flexibility Act.

List of Subjects in 14 CFR Part 71

Aviation safety, Transition areas.

Adoption of the Amendment

Accordingly, pursuant to the authority delegated to me, part 71 of the Federal Aviation Regulations (14 CFR part 71) is amended as follows:

PART 71—DESIGNATION OF FEDERAL AIRWAYS, AREA LOW ROUTES, CONTROLLED AIRSPACE, AND REPORTING POINTS

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

Authority: 49 U.S.C. 1348(a), 1354(a), 1510; Executive Order 10854; 49 U.S.C. 106(g) (Revised Pub. L. 97–449, January 12, 1983); 14 CFR 11.69.

§71.181 [Amended]

2. Section 71.181 is amended as follows:

McCall, Idaho [Revised]

That airspace extending upward from 700 feet above the surface within a 5-mile radius of McCall Municipal Airport (Latitude 44 53'23" N., Longitude 116 06'00" W.); that airspace extending upward from 1,200 feet above the surface within 6 miles west and 17 miles east of the McCall VORTAC 344 and 164 radials extending from 20 miles south to 19 miles north of the VORTAC.

Issued in Seattle, Washington, on December 11, 1989.

Temple H. Johnson, Jr.,

Manager, Air Traffic Division Northwest Mountain Region.

[FR Doc. 90–914 Filed 1–12–90; 8:45 am]

BILLING CODE 4910–13–M

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14 CFR Part 75

[Airspace Docket No. 89–ASO–16]

Alteration of Jet Route J–213; West Virginia

AGENCY: Federal Aviation Administration (FAA), DOT.

ACTION: Final rule.

SUMMARY: This amendment alters the description of Jet Route J–213, located in the vicinity of Beckley, WV, by extending that route from Beckley to Louisville, KY, via a south dogleg. The ARMEL standard terminal arrival (STAR) which began at Beckley for transition into several terminal areas is cancelled. The extension of J–213 improves the flow of traffic into several terminal areas and reduces controller workload.

EFFECTIVE DATE: 0901 u.t.c. March 8, 1990.

FOR FURTHER INFORMATION CONTACT: Lewis W. Still, Airspace Branch (ATO–240), Airspace-Rules and Aeronautical Information Division, Air Traffic Operations Service, Federal Aviation Administration, 800 Independence Avenue, SW., Washington, DC 20591; telephone: (202) 267–9250.

SUPPLEMENTARY INFORMATION:

History

On July 17, 1989, the FAA proposed to amend part 75 of the Federal Aviation Regulations (14 CFR part 75) to alter the description of Jet Route J–213 located in the vicinity of Beckley, WV, by extending that route from Beckley to Louisville, KY (54 FR 29909). The ARMEL STAR is established from Beckley as a transition route to several east coast airports. The ARMEL STAR is cancelled and is replaced by two STAR's using Beckley as the initial starting point. Extending J–213 via a south dogleg provides an established route in an area where aircraft are usually vectored. This action reduces controlled workload. Interested parties were invited to participate in this rulemaking proceeding by submitting written comments on the proposal to the FAA. No comments objecting to the proposal were received. Except for editorial changes, this amendment is the same as that proposed in the notice. Section 75.100 of part 75 of the Federal Aviation Regulations was republished in Handbook 7400.6E dated January 3, 1989.

The Rule

This amendment to part 75 of the Federal Aviation Regulations alters the description of Jet Route J–213, located in the vicinity of Beckley, WV, by extending that route from Beckley to Louisville, KY, via a south dogleg. The ARMEL STAR which begins at Beckley for transition into several terminal areas is cancelled. The extension of J–213 improves the flow of traffic into several terminal areas and reduces controller workload.

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. It, therefore—(1) is not a "major rule" under Executive Order 12291; (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 75

Aviation safety, Jet routes.

Adoption of the Amendment

Accordingly, pursuant to the authority delegated to me, part 75 of the Federal Aviation Regulations (14 CFR part 75) is amended, as follows:

PART 75—ESTABLISHMENT OF JET ROUTES AND AREA HIGH ROUTES

1. The authority citation for part 75 continues to read as follows:

Authority. 49 U.S.C. 1348(a), 1354(a), 1510; Executive Order 10854; 49 U.S.C. 106(g)

(Revised Pub. L. 97-449, January 12, 1983); 14 CFR 11.69.

§ 75.100 [Amended]

2. Section 75.100 is amended as follows:

J-213 [Amended]

By removing the words "to Beckley." and by substituting the words "Beckley; INT Beckley 264° and Louisville, KY, 101° radials; to Louisville."

Issued in Washington, DC, on January 4, 1990.

Harold W. Becker,

Manager, Airspace-Rules and Aeronautical Information Division.

[FR Doc. 90-915 Filed 1-12-90; 8:45 am]

BILLING CODE 4910-13-M

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

14 CFR Part 1221

NASA Seal and Other Devices, and the Congressional Space Medal of Honor

AGENCY: National Aeronautics and Space Administration (NASA).

ACTION: Final rule.

SUMMARY: NASA is amending 14 CFR part 1221 by revising one paragraph in subpart 1221.1, "NASA Seal, Insignia, Logotype Insignia, Program and Astronaut Badges, and Flags, and the Agency's Unified Visual Communications System." This revised paragraph adds the restricted use of a NASA Seal plaque in paragraph (a)(5) of § 1221.111.

EFFECTIVE DATE: January 16, 1990.

ADDRESSES: Public Services Division, NASA Headquarters, Washington, DC 20546.

FOR FURTHER INFORMATION CONTACT: Robert Schulman, (202) 453-8315.

SUPPLEMENTARY INFORMATION: Since this revision involves administrative and editorial management decisions and procedures, notice and public comment are not required.

The National Aeronautics and Space Administration has determined that:

1. This rule is not subject to the requirements of the Regulatory Flexibility Act, 5 U.S.C. 601-612, since it will not exert a significant economic impact on a substantial number of small entities.

2. This rule is not a major rule as defined in Executive Order 12291.

List of Subjects in 14 CFR Part 1221

Decorations, Medals, Awards, Flags, Seals, Insignia, Unified Visual Communications System.

For reasons set out in the Preamble, 14 CFR part 1221 is amended as follows:

PART 1221—THE NASA SEAL AND OTHER DEVICES, AND THE CONGRESSIONAL SPACE MEDAL OF HONOR

1. The authority citation for 14 CFR part 1221 continues to read as follows:

Authority: 42 U.S.C. 2472(a) and 2473(b)(1).

2. Section 1221.111 is amended by revising paragraph (a)(5) to read as follows:

§ 1221.111 Use of the NASA seal.

(a) * * *

(5) Plaques; the design of the NASA Seal may be incorporated in plaques for display in agency auditoriums, presentation rooms, lobbies, offices of senior officials, and on the fronts of buildings occupied by NASA. A separate NASA Seal in the form of a 15-inch, round, bronze colored plaque on a walnut colored wood base is also available, but prohibited for use in the above representational manners. It is restricted to use only as a presentation item by the Administrator and the Deputy Administrator.

* * * * *

Dated: January 8, 1990.

Richard H. Truly,

Administrator.

[FR Doc. 90-926 Filed 1-12-90; 8:45 am]

BILLING CODE 7510-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 14, 19, and 20

Public Hearing Before a Public Advisory Committee; Standards of Conduct and Conflicts of Interest; Public Information; Editorial Amendments

AGENCY: Food and Drug Administration.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending certain of its regulations for public hearings before a public advisory committee, standards of conduct and conflicts of interest, and public information to correct cross-references and to update a title. This action will improve the accuracy of the regulations.

EFFECTIVE DATE: January 16, 1990.

FOR FURTHER INFORMATION CONTACT: T. Rada Proehl, Regulations Editorial Staff (HFC-222), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-443-2994.

SUPPLEMENTARY INFORMATION: FDA is amending certain of its regulations for a public hearing before a public advisory committee, standards of conduct and conflicts of interest, and public information to correct cross-references and update a title. The amendments in 21 CFR 14.7(b), 19.10(a), 20.41(b)(4), and 20.47(c) are wholly editorial in nature. For this reason, FDA finds for good cause that notice and public procedure and delayed effective date are unnecessary (5 U.S.C. 553(b)(B) and (d)).

List of Subjects

21 CFR Part 14

Administrative practice and procedure, Advisory committees, Color additives, Drugs, Radiation protection.

21 CFR Part 19

Conflict of interests.

21 CFR Part 20

Confidential business information, Courts, Freedom of information, Government employees.

Therefore, under the Federal Advisory Committee Act, the Federal Food, Drug, and Cosmetic Act, and the Freedom of Information Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR parts 14, 19, and 20 are amended as follows:

PART 14—PUBLIC HEARING BEFORE A PUBLIC ADVISORY COMMITTEE

1. The authority citation for 21 CFR part 14 continues to read as follows:

Authority: Secs. 201-902 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321-392); 21 U.S.C. 41-50, 141-149, 467f, 679, 821, 1034; secs. 2, 351, 354-360F, 361 of the Public Health Service Act (42 U.S.C. 201, 262, 263b-263n, 264); secs. 2-12 of the Fair Packaging and Labeling Act (15 U.S.C. 1451-1461); 5 U.S.C. App. 2; 28 U.S.C. 2112.

§ 14.7 [Amended]

2. Section 14.7 *Administrative remedies* is amended in paragraph (b) by removing "45 CFR 5.82" and replacing it with "45 CFR 5.34".

PART 19—STANDARDS OF CONDUCT AND CONFLICTS OF INTEREST

3. The authority citation for 21 CFR part 19 continues to read as follows:

Authority: Sec. 701 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 371).

§ 19.10 [Amended]

4. Section 19.10 *Food and Drug Administration Conflict of Interest Review Board* is amended in paragraph

(a) by removing "Division of Human Resources Management" and replacing it with "Division of Ethics and Program Integrity".

PART 20—PUBLIC INFORMATION

5. The authority citation for 21 CFR part 20 continues to read as follows:

Authority: Secs. 201–902 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321–392); secs. 301, 302, 303, 307, 310, 311, 351, 352, 354–360F, 361, 362, 1701–1706, 2101 of the Public Health Service Act (42 U.S.C. 241, 242, 242a, 242l, 242n, 243, 262, 263, 263b–263n, 264, 265, 300u–300u–5, 300aa–1); 5 U.S.C. 552; 18 U.S.C. 1905.

§ 20.41 [Amended]

6. Section 20.41 *Time limitations* is amended in paragraph (b)(4) by removing "45 CFR 5.82" and replacing it with "45 CFR 5.34".

§ 20.47 [Amended]

7. Section 20.47 *Denial of a request for records* is amended in paragraph (c) by removing "45 CFR 5.82" and replacing it with "45 CFR 5.34".

Dated: January 5, 1990.

Ronald G. Chesemore,

Associate Commissioner for Regulatory Affairs.

[FR Doc. 90–933 Filed 1–12–90; 8:45 am]

BILLING CODE 4160–01–M

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21 CFR Parts 510 and 522

Animal Drugs, Feeds, and Related Products; Serum Gonadotropin and Chorionic Gonadotropin for Injection

AGENCY: Food and Drug Administration.

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 Intervet America, Inc. The NADA provides for subcutaneous use of reconstituted serum gonadotropin and chorionic gonadotropin for inducing fertile estrus (heat) in prepuberal (noncycling) gilts over 5½ months old and weighing at least 85 kilograms (kg) (185 pounds (lb)).

EFFECTIVE DATE: January 16, 1990.

FOR FURTHER INFORMATION CONTACT: James F. McCormack, Center for Veterinary Medicine (HFV–128), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–443–4317.

SUPPLEMENTARY INFORMATION: Intervet America, Inc., P.O. Box 318, 405 State St., Millsboro, DE 19966, filed NADA 140–856 providing for use of a freeze-dried serum gonadotropin and chorionic gonadotropin to be reconstituted for subcutaneous use for induction of fertile estrus (heat) in prepuberal (noncycling) gilts over 5½ months old and weighing at least 85 kg (185 lb). The NADA is approved as of January 5, 1990, and the regulations are amended by adding new § 522.1079 to reflect the approval. The basis for approval is discussed in the freedom of information summary.

In addition, Intervet America, Inc., has not been listed in the list of sponsors of approved applications. That list in § 510.600(c) (1) and (2) is amended to reflect the new sponsor.

In accordance with the freedom of information provisions of Part 20 (21 CFR Part 20) and § 514.11(e)(2)(ii) (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, Rm. 4–62, 5600 Fishers Lane, Rockville, MD 20857, from 9 a.m. to 4 p.m., Monday through Friday.

The agency has carefully considered the potential environmental effects of this action. FDA has concluded that the action will not have a significant impact on the human environment, and that an environmental impact statement is not required. The agency's finding of no significant impact and the evidence supporting that finding, contained in an environmental assessment, may be seen in the Dockets Management Branch (address above) between 9 a.m. and 4 p.m., Monday through Friday.

List of Subjects

21 CFR Part 510

Administrative practice and procedure, Animal drugs, Labeling, Reporting and recordkeeping requirements.

21 CFR Part 522

Animal drugs.

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 parts 510 and 522 are amended as follows:

PART 510—NEW ANIMAL DRUGS

1. The authority citation for 21 CFR part 510 continues to read as follows:

Authority: Secs. 201, 301, 501, 502, 503, 512, 701, 706 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 331, 351, 352, 353, 360b, 371, 376).

2. Section 510.600 is amended in the table in paragraph (c)(1) by alphabetically adding a new entry "Intervet America, Inc.", and in the table in paragraph (c)(2) by numerically adding a new entry "057926", to read as follows:

§ 510.600 Names, addresses, and drug labeler codes of sponsors of approved applications.

* * * * *

(c) * * *

(1) * * *

Firm name and address	Drug labeler code
* * * * *	
Intervet America, Inc., P.O. Box 318, 405 State St., Millsboro, DE 19966....	057926
* * * * *	

(2) * * *

Drug labeler code	Firm name and address
* * * * *	
057926.....	Intervet America, Inc., P.O. Box 318 405 State St., Millsboro, DE 19966

PART 522—IMPLANTATION OR INJECTABLE DOSAGE FORM NEW ANIMAL DRUGS NOT SUBJECT TO CERTIFICATION

3. The authority citation for 21 CFR part 522 continues to read as follows:

Authority: Sec. 512 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b).

4. New § 522.1079 is added to read as follows:

§ 522.1079 Serum gonadotropin and chorionic gonadotropin.

(a) *Specifications.* Each dose consists of 400 international units (I.U.) serum gonadotropin and 200 I.U. chorionic gonadotropin as a freeze-dried powder to be reconstituted with 5 milliliters of sterile aqueous diluent.

(b) *Sponsor.* See No. 057926 in § 510.600(c) of this chapter.

(c) *Conditions of use in swine.* (1) *Amount.* 400 I.U. serum gonadotropin with 200 I.U. chorionic gonadotropin per 5 milliliters dose per animal.

(2) *Indications for use.* For induction of fertile estrus (heat) in prepuberal (noncycling) gilts.

(3) *Limitations.* For subcutaneous use in prepuberal gilts over 5½ months old and weighing at least 85 kilograms (185 pounds).