ability to use trim switches to override uncommanded movement or yoke disconnect switches to disconnect the HSTA, which could result in reduction of or loss of pitch trim control and consequent reduced controllability of the airplane.

Compliance

(e) You are responsible for having the actions required by this AD performed within the compliance times specified, unless the actions have already been done.

Airplane Flight Manual (AFM) Revision

(f) Within 7 days after the effective date of this AD, make the applicable AFM revisions specified in paragraph (f)(1) or (f)(2) of this AD by incorporating the applicable Canadair (Bombardier) temporary revisions (TRs) identified in Table 1 of this AD into the applicable AFM.

(1) For Model CL–600–2B16 (CL–604) airplanes: Revise the Emergency Procedures section of the AFM to advise the flightcrew of additional procedures to follow in the event of stabilizer trim runaway.

(2) For Model CL–600–2B19 (Regional Jet Series 100 & 440) airplanes: Revise the Emergency and Abnormal Procedures sections of the AFM to advise the flightcrew of additional procedures to follow in the event of stabilizer trim runaway and in the event of MACH TRIM, STAB TRIM, and horizontal stabilizer trim malfunctions.

TABLE 1.—TRS

For Bombardier Model—	Use—	Dated—	To the—
CL–600–2B16 (CL–604) airplanes	Canadair Challenger TR 604/21.	August 1, 2006	Canadair Challenger CL–604 AFM, PSP 604–1.
CL–600–2B19 (Regional Jet Series 100 & 440) airplanes.	Canadair Regional Jet TR RJ/152–4.	August 9, 2006	Canadair Regional Jet AFM, CSP A–012.

(g) When the applicable TR has been included in the general revisions of the applicable AFM, those general revisions may be inserted into the AFM and the applicable TR may be removed, provided the relevant information in the general revisions is identical to that in the TR.

Alternative Methods of Compliance (AMOCs)

(h)(1) The Manager, New York Aircraft Certification Office, FAA, has the authority to approve AMOCs for this AD, if requested in accordance with the procedures found in 14 CFR 39.19.

(2) Before using any AMOC approved in accordance with § 39.19 on any airplane to which the AMOC applies, notify the appropriate principal inspector in the FAA Flight Standards Certificate Holding District Office.

Related Information

(i) Canadian airworthiness directives CF–2006–20, dated August 22, 2006, and CF–2006–21, dated August 23, 2006, also address the subject of this AD.

Material Incorporated by Reference

(j) You must use the applicable service information specified in Table 2 of this AD to perform the actions that are required by this AD, unless the AD specifies otherwise. The Director of the Federal Register approved the incorporation by reference of these documents in accordance with 5 U.S.C. 552(a) and 1 CFR part 51. Contact Bombardier, Inc., Canadair, Aerospace Group, P.O. Box 6087, Station Centre-ville, Montreal, Quebec H3C 3G9, Canada, for a copy of this service information. You may review copies at the Docket Management Facility, U.S. Department of Transportation, 400 Seventh Street SW., Room PL–401, Nassif Building, Washington, DC; on the Internet at *http://dms.dot.gov*; or at the National Archives and Records Administration (NARA). For information on the availability of this material at the NARA, call (202) 741–6030, or go to *http://www.archives.gov/federal_register/code_of_federal_regulations/ibr_locations.html*.

TABLE 2.—TEMPORARY REVISIONS INCORPORATED BY REFERENCE

Temporary revision	Dated—	To the—
Canadair Challenger Temporary Revision 604/21	August 1, 2006	Canadair Challenger CL–604 Airplane Flight Manual, PSP 604–1.
Canadair Regional Jet Temporary Revision RJ/152–4	August 9, 2006	Canadair Regional Jet Airplane Flight Manual, CSP A–012.

Issued in Renton, Washington, on August 29, 2006.

Ali Bahrami,

Manager, Transport Airplane Directorate, Aircraft Certification Service.

[FR Doc. E6–14617 Filed 8–31–06; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 71

[Docket No. 29334; Amendment No. 71–38]

Airspace Designations; Incorporation by Reference

AGENCY: Federal Aviation Administration (FAA), DOT.

ACTION: Final rule.

SUMMARY: This action amends Title 14 Code of Federal Regulations (14 CFR) part 71 relating to airspace designations to reflect the approval by the Director of the **Federal Register** of the incorporation by reference of FAA Order 7400.9P, Airspace Designations and Reporting Points. This action also explains the procedures the FAA will use to amend the listings of Class A, B, C, D, and E airspace areas; air traffic service routes; and reporting points incorporated by reference.

DATES: *Effective Date:* These regulations are effective September 15, 2006, through September 15, 2007. The incorporation by reference of FAA Order 7400.9P is approved by the Director of the Federal Register as of September 15, 2006, through September 15, 2007.

FOR FURTHER INFORMATION CONTACT: Tameka Bentley, Airspace and Rules, Office of System Operations Airspace

and AIM, Federal Aviation Administration, 800 Independence Avenue, SW., Washington, DC 20591; telephone: (202) 267–8783.

SUPPLEMENTARY INFORMATION:

History

FAA Order 7400.9N, Airspace Designations and Reporting Points, dated September 1, 2005, and effective September 15, 2005, listed Class A, B, C, D and E airspace areas; air traffic service routes; and reporting points. Due to the length of these descriptions, the FAA requested approval from the Office of the Federal Register to incorporate the material by reference in the Federal Aviation Regulations § 71.1, effective September 15, 2005, through September 15, 2006. During the incorporation by reference period, the FAA processed all proposed changes of the airspace listings in FAA Order 7400.9N in full text as proposed rule documents in the **Federal Register**. Likewise, all amendments of these listings were published in full text as final rules in the **Federal Register**. This rule reflects the periodic integration of these final rule amendments into a revised edition of Order 7400.9P, Airspace Designations and Reporting Points. The Director of the Federal Register has approved the incorporation by reference of FAA Order 7400.9P in § 71.1, as of September 15, 2006, through September 15, 2007. This rule also explains the procedures the FAA will use to amend the airspace designations incorporated by reference in part 71. Sections 71.5, 71.15, 71.31, 71.33, 71.41, 71.51, 71.61, 71.71, and 71.901 are also updated to reflect the incorporation by reference of FAA Order 7400.9P.

The Rule

This action amends Title 14 Code of Federal Regulations (14 CFR) part 71 to reflect the approval by the Director of the Federal Register of the incorporation by reference of FAA Order 7400.9P, effective September 15, 2006, through September 15, 2007. During the incorporation by reference period, the FAA will continue to process all proposed changes of the airspace listings in FAA Order 7400.9P in full text as proposed rule documents in the **Federal Register**. Likewise, all amendments of these listings will be published in full text as final rules in the **Federal Register**. The FAA will periodically integrate all final rule amendments into a revised edition of the order, and submit the revised edition to the Director of the Federal Register for approval for incorporation by reference in section 71.1.

The FAA has determined that this action: (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) does not warrant preparation of a regulatory evaluation as the anticipated impact is so minimal. This action neither places any new restrictions or requirements on the public, nor changes the dimensions or operation requirements of the airspace listings incorporated by reference in part 71. Consequently, notice and public procedure under 5 U.S.C. 553(b) are unnecessary. Because this action will continue to update the changes to the airspace designations, which are depicted on aeronautical charts, and to avoid any unnecessary pilot confusion, I find that good cause exists, under 5 U.S.C. 553(d), for making this amendment effective in less than 30 days.

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, B, C, D, AND E AIRSPACE AREAS; AIR TRAFFIC SERVICE ROUTES; AND REPORTING POINTS

■ 1. The authority citation for 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.

■ 2. Section 71.1 is revised to read as follows:

§ 71.1 Applicability.

The complete listing for all Class A, B, C, D, and E airspace areas; air traffic service routes; and reporting points can be found in FAA Order 7400.9P, Airspace Designations and Reporting Points, dated September 1, 2006. This incorporation by reference was approved by the Director of the Federal Register in accordance with 5 U.S.C. 552 (a) and 1 CFR part 51. The approval to incorporate by reference FAA Order 7400.9P is effective September 15, 2006, through September 15, 2007. During the incorporation by reference period, proposed changes to the listings of Class A, B, C, D, and E airspace areas; air traffic service routes; and reporting points will be published in full text as proposed rule documents in the **Federal Register**. Amendments to the listings of Class A, B, C, D, and E airspace areas; air traffic service routes; and reporting points will be published in full text as final rules in the **Federal Register**. Periodically, the final rule amendments will be integrated into a revised edition of the Order and submitted to the Director of the Federal Register for approval for incorporation by reference in this section. Copies of FAA Order 7400.9P may be obtained from the Law Library, Office of the Chief Counsel, Federal Aviation Administration, 800 Independence Avenue, SW., Washington, DC 20591, (202) 267–3174. An electronic version of the Order is available on the FAA Web site at *http://www.faa.gov/ATPUBS*. Copies of FAA Order 7400.9P may be inspected in Docket No. 29334, at the same address above, weekdays between 8 a.m. and 4:30 p.m., or on the **Federal Register** Web site at *www.regulations.gov*.

§ 71.5 [Amended]

■ 3A. Section 71.5 is amended by removing the words ''FAA Order 7400.9N'' and adding,in their place, the words ''FAA Order 7400.9P.''

§ 71.15 [Amended]

■ 3B. Section 71.5 is amended by removing the words ''FAA Order 7400.9N'' and adding, in their place, the words ''FAA Order 7400.9P.''

§ 71.31 [Amended]

■ 4. Section 71.31 is amended by removing the words ''FAA Order 7400.9N'' and adding, in their place, the words ''FAA Order 7400.9P.''

§ 71.33 [Amended]

■ 5. Paragraph (c) of section 71.33 is amended by removing the words ''FAA Order 7400.9N'' and adding, in their place, the words ''FAA Order 7400.9P.''

§ 71.41 [Amended]

■ 6. Section 71.41 is amended by removing the words ''FAA Order 7400.9N'' and adding, in their place, the words ''FAA Order 7400.9P.''

§ 71.51 [Amended]

■ 7. Section 71.51 is amended by removing the words ''FAA Order 7400.9N'' and adding, in their place, the words ''FAA Order 7400.9P.''

§ 71.61 [Amended]

■ 8. Section 71.61 is amended by removing the words ''FAA Order 7400.9N'' and adding, in their place, the words ''FAA Order 7400.9P.''

§ 71.71 [Amended]

■ 9. Paragraph (b), (c), (d), (e), and (f) of section 71.71 are amended by removing

the words ''FAA Order 7400.9N'' and adding, in their place, the words ''FAA Order 7400.9P.''

§ 71.901 [Amended]

■ 10. Paragraph (a) of section 71.901 is amended by removing the words ''FAA Order 7400.9N'' and adding, in their place, the words ''FAA Order 7400.9P.''

Issued in Washington, DC, on July 27, 2006.

Edith V. Parish,

Manager, Airspace and Rules.

[FR Doc. E6–12434 Filed 8–31–06; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 520

Oral Dosage Form New Animal Drugs; Carprofen

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 supplemental abbreviated new animal drug application (ANADA) filed by IMPAX Laboratories, Inc. The supplemental ANADA provides for veterinary prescription use of carprofen caplets in dogs for the control of postoperative pain associated with soft tissue and orthopedic surgeries.

DATES: This rule is effective September 1, 2006.

FOR FURTHER INFORMATION CONTACT: John K. Harshman, Center for Veterinary Medicine (HFV–104), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 240–276–9808, e-mail: *john.harshman@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: IMPAX Laboratories, Inc., 30831 Huntwood Ave., Hayward, CA 94544, filed a supplement to ANADA 200–366 for NOVOX (carprofen) caplets which are approved for veterinary prescription use in dogs for the relief of pain and inflammation associated with osteoarthritis (70 FR 30625, May 27, 2005). The supplement provides for use of NOVOX caplets for the control of postoperative pain associated with soft tissue and orthopedic surgeries. The supplemental ANADA is approved as of July 27, 2006, and 21 CFR 520.309 is amended to reflect the approval.

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 Division of Dockets Management (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.

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 Subjects in 21 CFR Part 520

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 part 520 is amended as follows:

PART 520—ORAL DOSAGE FORM NEW ANIMAL DRUGS

■ 1. The authority citation for 21 CFR part 520 continues to read as follows:

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

■ 2. In § 520.309, remove paragraphs (d)(2)(i) and (d)(2)(ii), and revise paragraphs (b)(2) and (d)(2) to read as follows:

§ 520.309 Carprofen.

* * * * *

(b) * * *

(2) No. 000115 for use of product described in paragraph (a)(1) as in paragraph (d) of this section.

* * * * *

(d) * * *

(2) *Indications for use.* For the relief of pain and inflammation associated with osteoarthritis and for the control of postoperative pain associated with soft tissue and orthopedic surgeries.

* * * * *

Dated: August 18, 2006.

Steven D. Vaughn,

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

[FR Doc. E6–14508 Filed 8–31–06; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 522

Implantation or Injectable Dosage Form New Animal Drugs; Lincomycin

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 an abbreviated new animal drug application (ANADA) filed by Cross Vetpharm Group Ltd. The ANADA provides for the use of lincomycin injectable solution in swine for the treatment of infectious arthritis and mycoplasma pneumonia.

DATES: This rule is effective September 1, 2006.

FOR FURTHER INFORMATION CONTACT: John K. Harshman, Center for Veterinary Medicine (HFV–104), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0169, e-mail: *john.harshman@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: Cross Vetpharm Group Ltd., Broomhill Rd., Tallaght, Dublin 24, Ireland, filed ANADA 200–368 that provides for use of LINCOMED 100 (lincomycin hydrochloride) and LINCOMED 300 (lincomycin hydrochloride) in swine for the treatment of infectious arthritis and mycoplasma pneumonia. Cross Vetpharm Group Ltd.'s LINCOMED 100 and LINCOMED 300 are approved as generic copies of LINCOMIX 100 Injectable and LINCOMIX 300 Injectable, sponsored by Pharmacia & Upjohn Co., a Division of Pfizer, Inc., under NADA 034 025. The ANADA is approved as of July 27, 2006, and the regulations are amended in § 522.1260 to reflect the approval. The basis of approval is discussed in the freedom of information summary.

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 Division of Dockets Management (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.

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