contain aircraft executing Standard Instrument Approach Procedures (SIAPs). Two new Standard Instrument Approach Procedures (SIAPs) are being developed for the New Stuyahok Airport. This action revises existing Class E airspace upward from 700 feet (ft.) and 1,200 ft. above the surface at New Stuyahok Airport, New Stuyahok, AK.

EFFECTIVE DATE: 0901 UTC, June 5, 2008. The Director of the Federal Register approves this incorporation by reference action under title 1, Code of Federal Regulations, part 51, subject to the annual revision of FAA Order 7400.9 and publication of conforming amendments.

FOR FURTHER INFORMATION CONTACT: Gary Rolf, AAL–538G, Federal Aviation Administration, 222 West 7th Avenue, Box 14, Anchorage, AK 99513–7587; telephone number (907) 271–5898; fax: (907) 271–2850; e-mail: *gary.ctr.rolf@faa.gov*. Internet address: *http://www.alaska.faa.gov/at*.

SUPPLEMENTARY INFORMATION:

History

On Friday, February 1, 2008, the FAA proposed to amend part 71 of the Federal Aviation Regulations (14 CFR part 71) to revise Class E airspace upward from 700 ft. above the surface and from 1,200 ft. above the surface at New Stuyahok, AK (73 FR 6057). The action was proposed in order to create Class E airspace sufficient in size to contain aircraft while executing SIAPs for the New Stuyahok Airport. The Notice of Proposed Rulemaking contained airport location data, which has since been updated. The revised airport location coordinates are listed in this rule. Class E controlled airspace extending upward from 700 ft. above the surface and from 1,200 ft. above the surface in the New Stuyahok Airport area is revised by this action.

Interested parties were invited to participate in this rulemaking proceeding by submitting written comments on the proposal to the FAA. No comments were received. The rule is adopted as proposed.

The area will be depicted on aeronautical charts for pilot reference. The coordinates for this airspace docket are based on North American Datum 83. The Class E airspace areas designated as 700/1,200 ft. transition areas are published in paragraph 6005 of FAA Order 7400.9R, *Airspace Designations and Reporting Points*, signed August 15, 2007, and effective September 15, 2007, which is incorporated by reference in 14 CFR 71.1. The Class E airspace designations listed in this document will be published subsequently in the Order.

The Rule

This amendment to 14 CFR part 71 revises Class E airspace at the New Stuyahok Airport, Alaska. This Class E airspace is revised to accommodate aircraft executing new SIAPs, and will be depicted on aeronautical charts for pilot reference. The intended effect of this rule is to provide adequate controlled airspace for Instrument Flight Rules (IFR) operations at the New Stuyahok Airport, New Stuyahok, Alaska.

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 ''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. 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.

The FAA's authority to issue rules regarding aviation safety is found in Title 49 of the United States Code. Subtitle 1, Section 106 describes the authority of the FAA Administrator. Subtitle VII, Aviation Programs, describes in more detail the scope of the agency's authority.

This rulemaking is promulgated under the authority described in Subtitle VII, Part A, Subpart 1, Section 40103, Sovereignty and use of airspace. Under that section, the FAA is charged with prescribing regulations to ensure the safe and efficient use of the navigable airspace. This regulation is within the scope of that authority because it creates Class E airspace sufficient in size to contain aircraft executing instrument procedures for the New Stuyahok Airport and represents the FAA's continuing effort to safely and efficiently use the navigable airspace.

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 7400.9R, *Airspace Designations and Reporting Points*, signed August 15, 2007, and effective September 15, 2007, 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 New Stuyahok, AK [Revised]

New Stuyahok, New Stuyahok Airport, AK
(Lat. 59°27′06″ N., long. 157°22′23″ W.)

That airspace extending upward from 700 feet above the surface within a 6.9-mile radius of the New Stuyahok Airport; and that airspace extending upward from 1,200 feet above the surface within a 71-mile radius of the New Stuyahok Airport.

* * * * *

Issued in Anchorage, AK, on March 24, 2008.

Anthony M. Wylie,

Manager, Alaska Flight Services Information Area Group.

[FR Doc. E8–6921 Filed 4–3–08; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 210 and 211

[Docket No. FDA–2008–N–0179] (formerly Docket No. 2007N–0280)

Amendment to the Current Good Manufacturing Practice Regulations for Finished Pharmaceuticals; Withdrawal

AGENCY: Food and Drug Administration, HHS.

ACTION: Direct final rule; withdrawal.

SUMMARY: The Food and Drug Administration (FDA) is withdrawing a

direct final rule that published in the **Federal Register** of December 4, 2007 (72 FR 68064), to amend certain regulations as the first phase of an incremental approach to modernize or clarify some of the current good manufacturing practice (CGMP) regulations for finished pharmaceuticals, as well as harmonize some of the CGMP requirements with those of other foreign regulators and other FDA regulations. The comment period closed February 19, 2008. FDA is withdrawing the direct final rule because the agency received significant adverse comments. FDA will consider the comments received under our usual procedures for notice and comment in connection with the notice of proposed rulemaking that was published in the **Federal Register** of December 4, 2007, as a companion to the direct final rule (72 FR 68113).

DATES: The direct final rule published at 72 FR 68064 on December 4, 2007, is withdrawn as of April 4, 2008.

FOR FURTHER INFORMATION CONTACT:
Mary Malarkey, Center for Biologics Evaluation and Research (HFM–600), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852–1448, 301–827–6190, or
Dennis Bensley, Center for Veterinary Medicine (HFV–140), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 240–276–8268, or
Brian Hasselbalch, Center for Drug Evaluation and Research (HFD–320), Food and Drug Administration, 10903 New Hampshire Ave., Silver Spring, MD 20993, 301–796–3279.

Authority: Therefore, under the Federal Food, Drug, and Cosmetic Act, and under authority delegated to the Commissioner of Food and Drugs, the direct final rule published on December 4, 2007 (72 FR 68064) is withdrawn.

Dated: March 24, 2008.

Jeffrey Shuren,

Associate Commissioner for Policy and Planning.

[FR Doc. E8–7107 Filed 4–3–08; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510, 520, 526, and 558

Animal Drugs, Feeds, and Related Products; Withdrawal of Approval of NADAs; 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 by removing those portions that reflect approval of seven new animal drug applications (NADAs) because FDA is withdrawing approval of the NADAs.

DATES: This rule is effective April 4, 2008.

FOR FURTHER INFORMATION CONTACT: Pamela K. Esposito, Center for Veterinary Medicine (HFV–212), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–276–9067; e-mail: *pamela.esposito@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: The following sponsors have requested that FDA withdraw approval of the seven NADAs listed below because the products are no longer manufactured or marketed:

Sponsor	NADA Number Product (Drug)	21 CFR Cite Affected (Sponsor Drug Labeler Code)
Eon Labs Manufacturing, Inc., 227–15 North Conduit Ave., Laurelton, NY 11413	NADA 65–063, Tetracycline capsules	520.2345a (000185)
	NADA 65–345, Chloramphenicol capsules	520.390b (000185)
G.C. Hanford Manufacturing Co., P.O. Box 1017, Syracuse, NY 13201	NADA 65–465, AQUA-MAST (penicillin G procaine)	526.1696a (010515)
International Nutrition, Inc., 7706 "I" Plaza, Omaha, NE 68127	NADA 95–551, TYLAN 5 Premix (tylosin phosphate)	558.625 (043733)
	NADA 109–688, HYGROMIX 2.4 Premix (hygromycin B)	558.274 (043733)
	NADA 109–816, TYLAN 10 SULFA-G Premix (tylosin phosphate and sulfamethazine)	558.630 (043733)
Pfizer, Inc., 235 East 42d St., New York, NY 10017	NADA 103–758, TERAMIX–10 Premix (oxytetracycline)	Not codified

Following the withdrawal of approval of these NADAs, Eon Labs Manufacturing, Inc., is no longer sponsor of an approved application. Therefore, 21 CFR 510.600(c) is amended to remove entries for this sponsor.

As provided below, the animal drug regulations are amended to reflect the withdrawal of approvals. The regulations for penicillin G procaine