dated September 10, 1998, and effective September 16, 1998, is amended as follows:

Paragraph 6005 Class E Airspace Areas Extending Upward from 700 feet or More above the Surface of the Earth.

* * * * *

ASO GA E5 Carrollton, GA [Revised]

West Georgia Regional Airport
(Lat. 33°37′52″N, long. 85°09′07″W)
Carrolton NDB
(Lat. 33°33′57″N, long. 85°07′51″W)

That airspace extending upward from 700 feet or more above the surface of the earth within a 6.4-mile radius of West Georgia Regional Airport and within 3.5 miles from each side of the 166 degree bearing from the Carrollton NDB, extending from the 6.4-mile radius to 7 miles south of the NDB.

* * * * *

Issued in College Park, Georgia, on January 6, 1999.

Nancy B. Shelton,

Acting Manager, Air Traffic Division, Southern Region.

[FR Doc. 99–730 Filed 1–12–99; 8:45 am]

BILLING CODE 4910–13–M

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 71

[Airspace Docket No. 98–AEA–40]

Amendment to Class E Airspace; Romulus, NY

AGENCY: Federal Aviation Administration, (FAA) DOT.

ACTION: Final rule.

SUMMARY: This action removes Class E airspace at Seneca Army Air Field (AAF), Romulus, NY. The airport has been closed and all instrument procedures for the airport have been cancelled. The need for Class E airspace no longer exists for Instrument Flight Rules (IFR) operations at the airport. This action will result in the airspace reverting to Class E airspace.

EFFECTIVE DATE: 9091 UTC, March 25, 1999.

FOR FURTHER INFORMATION CONTACT: Mr. Francis Jordan, Airspace Specialist, Airspace Branch, AEA–520, Air Traffic Division, Eastern Region, Federal Aviation Administration, Federal #111, John F. Kennedy International Airport, Jamaica, New York 11430, telephone: (718) 553–4521.

SUPPLEMENTARY INFORMATION:

History

On November 3, 1998, a proposal to amend Part 71 of the Federal Aviation Regulations (14 CFR Part 71) to remove the Class E airspace extending upward from 700 feet above the surface at Seneca AAF, Romulus, NY, was published in the **Federal Register** (63 FR 59256).

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

The coordinates for this airspace docket are based on North American Datum 83. Class E airspace areas designations for airspace extending upward from 700 feet AGL are published in paragraph 6005 of FAA Order 7400.9F, dated September 10, 1998, and effective September 16, 1998, which is incorporated by reference in 14 CFR 71.1. The Class E airspace designation listed in this document will be removed subsequently from the Order.

The Rule

This amendment to Part 71 of the Federal Aviation Regulations (14 CFR Part 71) removes Class E airspace at Romulus, NY. The need for controlled airspace extending from 700 feet AGL at Seneca AAF no longer exists. This area will be removed from the appropriate aeronautical charts.

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 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 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—[AMENDED]

1. The authority citation for 14 CFR part 71 continues to read as follows:

Authority: 49 U.S.C. 106(g), 40103, 40113, 40120; EO 10854, 24 FR 9565, 3 CFR, 1959–1963 Comp., p. 389.

§ 71.1 [Amended]

The incorporation by reference in 14 CFR 71.1 of Federal Aviation Administration Order 7400.9F, Airspace Designations and Reporting Points, dated September 10, 1998, and effective September 16, 1998, is amended as follows:

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

* * * * *

AEA NY E5, Romulus, NY [Removed]

* * * * *

Issued in Jamaica, New York on January 4, 1999.

Franklin D. Hatfield,

Manager, Air Traffic Division, Eastern Region.

[FR Doc. 99–729 Filed 1–12–99; 8:45 am]

BILLING CODE 4910–13–M

SECURITIES AND EXCHANGE COMMISSION

17 CFR Part 279

[Release No. IA–1733A; File No. S7–28–97]

RIN 3235–AH22

Technical Changes to Schedule I to Form ADV

AGENCY: Securities and Exchange Commission.

ACTION: Final rule; technical changes to a form.

SUMMARY: The Commission is making technical changes to Schedule I to Form ADV, referenced in 17 CFR 279.1. Schedule I is the form on which investment advisers declare their eligibility for Commission registration. Schedule I to Form ADV was published Thursday, May 22, 1997 (62 FR 28112), under the Investment Advisers Act of 1940. Amendments to Schedule I to Form ADV were published Friday, July 24, 1998 (63 FR 39708), under the Advisers Act.

EFFECTIVE DATE: The rule amendments will become effective on January 7, 1999.

FOR FURTHER INFORMATION CONTACT: Arthur Laby, Special Counsel, at (202) 942–0716, Task Force on Investment Adviser Regulation, Division of Investment Management, Securities and Exchange Commission, 450 Fifth Street,

NW, Mail Stop 5–6, Washington, DC 20549.

I. Supplementary Information

Under section 203A of the Investment Advisers Act of 1940 (''Advisers Act''), the Commission has regulatory responsibility for an investment adviser that has at least $25 million of assets under management or advises a registered investment company. The Commission also has responsibility for an adviser that has less than $25 million of assets under management, if its principal office and place of business is in a state that has not enacted investment adviser legislation.[1] An adviser with its principal office in one of those states must indicate its eligibility for Commission registration on Schedule I of Form ADV.[2]

Colorado and Iowa recently passed investment adviser statutes, which became effective on January 1, 1999. An adviser that has its principal office and place of business in Colorado or Iowa, therefore, may not register with the Commission unless it has at least $25 million of assets under management, advises an investment company, or qualifies for an exemption under rule 203A–2.[3] Last July, the Commission adopted certain amendments to Schedule I to Form ADV.[4] The Commission today is making additional technical changes to Schedule I and the Instructions to Schedule I to reflect enactment of the Colorado and Iowa legislation.

New advisers (*i.e.*, those advisers that are not currently registered with the Commission) that have their principal place of business in Colorado or Iowa that are not eligible for Commission registration (*e.g.*, because they do not have at least $25 million of assets under management) must now register with Colorado or Iowa.[5] Advisers currently registered with the Commission solely because their principal office and place of business is located in Colorado or Iowa must withdraw from Commission registration no later than 180 days after the end of their fiscal year.[6]

II. Certain Findings

Under the Administrative Procedure Act (''APA''), notice of proposed rulemaking is not required when the agency, for good cause, finds ''that notice and public procedure thereon are impracticable, unnecessary, or contrary to the public interest.'' [7] The Commission is making technical changes to Schedule I to Form ADV to accommodate new legislation in Colorado and Iowa. The Commission, therefore, finds that publishing the changes for comment is unnecessary.

Publication of a substantive rule not less than 30 days before its effective date is required by the APA except as otherwise provided by the agency for good cause.[8] For the same reasons described above with respect to notice and opportunity for comment, the Commission finds that there is good cause for making these technical changes effective on January 7, 1999.

List of Subjects in 17 CFR Part 279

Reporting and recordkeeping requirements; Securities.

Accordingly, 17 CFR part 279 is amended as follows:

PART 279—FORMS PRESCRIBED UNDER THE INVESTMENT ADVISERS ACT OF 1940

1. The authority citation for part 279 continues to read as follows:

Authority: The Investment Advisers Act of 1940, 15 U.S.C. 80b–1, *et seq.*

2. By amending Schedule I to Form ADV (referenced in § 279.1) to remove all references to ''Colorado'' and ''Iowa'' and by amending the Instructions to Schedule I to Form ADV (referenced in § 279.1) to remove references to ''Colorado'' and ''Iowa'' and to remove the second paragraph under ''Instruction 3.''

Note: The text of Schedule I to Form ADV (§ 279.1) does not and the corrections will not appear in the Code of Federal Regulations.

Dated: January 7, 1999.

Margaret H. McFarland,

Deputy Secretary.

[FR Doc. 99–738 Filed 1–12–99; 8:45 am]

BILLING CODE 8010–01–U

[1] 15 U.S.C. 80b–3a.

[2] 17 CFR 279.1. Under rule 203–1 (17 CFR 275.203–1), an adviser must file Schedule I to Form ADV with its initial application for Commission registration, and under rule 204–1 (17 CFR 275.204–1), an adviser must file Schedule I to Form ADV with annual amendments to Form ADV.

[3] 17 CFR 275.203A–2.

[4] *See* Exemption for Investment Advisers Operating in Multiple States; Revisions to Rules Implementing Amendments to the Investment Advisers Act of 1940; Investment Advisers with Principal Offices and Places of Business in Colorado or Iowa, Investment Advisers Act Release No. 1733 (July 17, 1998) (63 FR 39708 (July 24, 1998)).

[5] In addition, advisers ineligible for Commission registration that have their principal office in Colorado or Iowa may be required to register in another state, if they have six or more clients that are residents of that state or have a place of business in that state. *See* Advisers Act section 222(d) (15 U.S.C. 80b–22(d)).

[6] Under rule 204–1(a) (17 CFR 275.204–1), an adviser is required to file its annual amendment to Form ADV within 90 days of the end of its fiscal year. Under rule 203A–1(c) (17 CFR 275.203A–1(c)), an adviser that is no longer eligible for Commission registration must withdraw from Commission registration within 90 days from the date the adviser was required to file its amended Form ADV. *See also* Schedule I to Form ADV, Instruction 6 (17 CFR 279.1).

[7] 5 U.S.C. 553(b).

[8] 5 U.S.C. 553(d).

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 520

Oral Dosage Form New Animal Drugs; Selegiline Hydrochloride Tablets

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 new animal drug application (NADA) filed by Pfizer, Inc. The supplemental NADA provides for oral veterinary prescription use of selegiline hydrochloride tablets for dogs for the control of clinical signs associated with cognitive dysfunction syndrome.

EFFECTIVE DATE: January 13, 1999.

FOR FURTHER INFORMATION CONTACT: Melanie R. Berson, Center For Veterinary Medicine (HFV–110), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–7543.

SUPPLEMENTARY INFORMATION: Pfizer, Inc., 235 East 42d St., New York, NY 10017, filed supplemental NADA 141–080 that provides for oral veterinary prescription use of Anipryl® (selegiline hydrochloride) tablets for dogs for the control of clinical signs associated with canine cognitive dysfunction syndrome. The product is approved for the control of clinical signs associated with uncomplicated pituitary-dependent hyperadrenocorticism. The supplement is approved as of December 10, 1998, and the regulations are amended by revising 21 CFR 520.2098 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 514.11(e)(2)(ii), a summary of safety and effectiveness data and information submitted to support approval of this supplemental application may be seen in the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, between 9 a.m. and 4 p.m., Monday