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DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 71

[Airspace Docket No. 99–AAL–4]

Amendment to Class E Airspace; Anaktuvuk Pass, AK

AGENCY: Federal Aviation Administration (FAA), DOT.

ACTION: Final rule, correction.

SUMMARY: This action corrects the error in the title of a correction to final rule that was published in the **Federal Register** on February 1, 1999 (64 FR 4784). The final rule establishing Class E airspace area at Anaktuvuk Pass, AK, was published in the **Federal Register** on November 5, 1998 (63 FR 59705), Airspace Docket 98–AAL–16.

EFFECTIVE DATE: 0901 UTC, July 16, 1999.

FOR FURTHER INFORMATION CONTACT: Robert van Haastert, Operations Branch, AAL–538, Federal Aviation Administration, 222 West 7th Avenue, Box 14, Anchorage, AK 99513–7587; telephone number (907) 271–5863; fax: (907) 271–2850; email: Robert.ctr.van-Haastert@faa. gov. Internet address: http://www.alaska.faa.gov/at or at address http://162.58.28.41/at.

SUPPLEMENTARY INFORMATION:

History

Federal Register Document 98–29627, Airspace Docket 98–AAL–16, published on November 5, 1998, (63 FR 59705) established the Class E airspace area at Anaktuvuk Pass, AK, **Federal Register** Document 99–2335, Airspace Docket 98–AAL–24, published February 1, 1999 (64 FR 4784) corrected an error in the geographic coordinates for the Anaktuvuk Pass Airport and Anaktuvuk Pass Non-Directional Radio Beacon. In the correction to final rule, Airspace Docket 98–AAL–24, the title for the Anaktuvuk Pass Class E airspace description is in error. The title ''AAL AK E2 Anaktuvuk Pass, AK'' should read ''AAL AK E5 Anaktuvuk Pass, AK''. This action corrects that error.

Correction to Final Rule

Accordingly, pursuant to the authority delegated to me, the title listed for the Anaktuvuk Pass airspace as published in the **Federal Register** on February 1, 1999 (64 FR 4784), (**Federal Register** Document 99–2335, page 4785), is corrected as follows:

§ 71.1 [Corrected]

* * * * *

AAL AK E5 Anaktuvuk Pass, AK [Corrected]

By removing ''AAL AK E2 Anaktuvuk Pass, AK'' and replacing with ''AAL AK E5 Anaktuvuk Pass, AK''.

* * * * *

Issued in Anchorage, AK, on June 3, 1999.

Trent S. Cummings,

Assistant Manager, Air Traffic Division, Alaskan Region.

[FR Doc. 99–15295 Filed 6–15–99; 8:45 am]

BILLING CODE 4910–13–M

FEDERAL TRADE COMMISSION

16 CFR Part 4

Miscellaneous Rules: Disclosure Requests

AGENCY: Federal Trade Commission.

ACTION: Final rule.

SUMMARY: The Commission is amending a rule of practice and procedure that governs disclosure requests. These amendments add requests for voluntary testimony to the scope of the rule's coverage. The amendments also clarify the existing scope of various paragraphs of the rule.

EFFECTIVE DATE: June 16, 1999.

FOR FURTHER INFORMATION CONTACT: Gary M. Greenfield, (202) 326–2753, Office of the General Counsel, Federal Trade Commission, 600 Pennsylvania Ave., NW, Washington, DC 20580.

SUPPLEMENTARY INFORMATION: The Commission is amending 16 CFR 4.11(e), which governs compulsory process requiring disclosure by Commission employees of material and information relating to their official duties. This provision also governs compulsory process to former Commission employees and to current and former special government employees that requires the disclosure of nonpublic information acquired during their Commission employment.

The amendments expand the scope of § 4.11(e) to include requests for voluntary testimony. As with requests by compulsory process for documents or testimony, the amended Rule requires anyone seeking voluntary testimony from Commission employees (and, where applicable, special government employees or former employees) to furnish a statement to the General Counsel setting forth information that will enable the General Counsel to make an informed decision regarding the request.

Amendments to paragraphs (c) and (d) of § 4.11 clarify that paragraph (e) of that section governs compulsory process from government agencies for Commission documents or testimony. Paragraph (e)(3), as amended, provides that the General Counsel may discretionarily waive the statement required by the Rule with respect to any individual request by a government agency.

The requirements of § 4.11(e) do not apply to invitations to testify before Congress or to testify before other government bodies on the possible effects of proposed legislation or regulations.

The Commission does not seek public comment on these amendments because they relate solely to agency practice and procedure. Thus, the amendments are exempt from the notice-and-comment requirements of the Administrative Procedure Act. *See* 5 U.S.C. 553(b)(A). In addition, the Commission certifies that these amendments will not have a significant impact on small business entities. Accordingly, no final regulatory flexibility analysis is required by the Regulatory Flexibility Act. *See* 5 U.S.C. 605(b).

List of Subjects in 16 CFR Part 4

Administrative practice and procedure.

For the reasons set forth in the preamble, the Commission amends part 4 of 16 CFR as follows:

PART 4—MISCELLANEOUS RULES

1. The authority for part 4 continues to read as follows:

Authority: Sec. 6, 38 Stat. 721; 15 U.S.C. 46.

2. Section 4.11 is amended by adding a sentence at the end of paragraphs (c) and (d) and revising paragraph (e) to read as follows:

§ 4.11 Disclosure requests.

* * * * *

(c) * * * Requests for material pursuant to compulsory process, or for voluntary testimony, in cases or matters in which the Commission is not a party will be treated in accordance with paragraph (e) of this section.

(d) * * * Request for material pursuant to compulsory process, or for voluntary testimony, in cases or matters in which the Commission is not a party will be treated in accordance with paragraph (e) of this section.

(e) *Requests for testimony, pursuant to compulsory process or otherwise, and requests for material pursuant to compulsory process, in cases or matters to which the Commission is not a party.* (1) The procedures specified in this section will apply to compulsory process and requests for voluntary testimony directed to Commission employees, except special government employees, that relate in any way to the employees' official duties. These procedures will also apply to compulsory process and requests for voluntary testimony directed to former Commission employees or to current or former special government employees of the Commission that seek nonpublic materials or information acquired during Commission employment. The provisions of paragraph (e)(3) of this section will also apply when requests described above are directed to the Commission. For purposes of this section, the term *testimony* includes any written or oral statement by a witness, such as depositions, affidavits, declarations, and statements at a hearing or trial; the term *nonpublic* includes any material or information which, under § 4.10, is not required to be more public; the term *employees,* except where otherwise specified, includes *special government employees* and other Commission employees; and the term *special government employees* includes consultants and other employees as defined by section 202 of title 18 of the United States Code.

(2) Any employee or former employee who is served with compulsory process shall promptly advise the General Counsel of its service, the nature of the material or information sought, and all relevant facts and circumstances. This notification requirement also applies to any employee or former employee whose testimony is sought on a voluntary basis under the conditions set forth in paragraph (e)(1) of this section.

(3) A party who causes compulsory process to be issued to, or who requests testimony by, the Commission or any employee or former employee of the Commission shall furnish a statement to the General Counsel, unless, with respect to a request by a Federal or State agency, the General Counsel determines, as a matter of discretion, to waive this requirement. The statement shall set forth the party's interest in the case or matter, the relevance of the desired testimony or material, and a discussion of whether it is reasonably available from other sources. If testimony is desired, the statement shall also contain a general summary of the testimony and a discussion of whether Commission records could be produced and used in its place. Any authorization for testimony will be limited to the scope of the demand as summarized in such statement.

(4) Absent authorization from the General Counsel, the employee or former employee shall respectfully decline to produce requested material or to disclose requested information. The refusal should be based on this paragraph and on *United States ex rel. Touhy* v. *Ragen,* 340 U.S. 462 (1951).

(5) The General Counsel will consider and act upon compulsory process and requests for voluntary testimony under this section with due regard for statutory restrictions, the Commission's rules and the public interest, taking into account such factors as the need to conserve the time of employees for conducting official business; the need to avoid spending the time and money of the United States for private purposes; the need to maintain impartiality between private litigants in cases where a substantial government interest is not involved; and the established legal standards for determining whether justification exists for the disclosure of confidential information and material.

(6) Invitations to testify before Congressional committees or subcommittees or to testify before other government bodies on the possible effects of legislative and regulatory proposals are not subject to paragraphs (e)(1) through (5) of this section.

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By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 99–15187 Filed 6–15–99; 8:45 am]

BILLING CODE 6750–01–M

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 new animal drug application (NADA) filed by Pfizer, Inc. The NADA provides for veterinary prescription use of carprofen chewable tablets for the relief of pain and inflammation associated with osteoarthritis in dogs.

EFFECTIVE DATE: June 16, 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 10017–5755, filed NADA 141–111 that provides for oral veterinary prescription use of Rimadyl® (carprofen) chewable tablets for the relief of pain and inflammation associated with osteoarthritis in dogs. The NADA is approved as of May 14, 1999. The regulations are amended in 21 CFR 520.309 by revising the section heading, by revising paragraph (a), by redesignating paragraph (c) as paragraph (d), by reserving paragraph (c), and by revising newly redesignated paragraphs (d)(1) and (d)(2) to reflect the approval.

The regulations currently provide for use of carprofen caplets in NADA 141–053. A revision of the indications for use has been approved by letter of April 21, 1999. At this time, the regulation is amended to reflect that approval.

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 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 through Friday.

Under section 512(c)(2)(F)(ii) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b(c)(2)(F)(ii)), this approval for nonfood-producing animals qualifies for 3 years of marketing