An informal docket may also be examined during normal business hours at the office of the Regional Air Traffic Division.

FOR FURTHER INFORMATION CONTACT: Patricia P. Crawford, Airspace and Obstruction Evaluation Branch (ATP–240), Airspace-Rules and Aeronautical Information Division, Air Traffic Rules and Procedures Service, Federal Aviation Administration, 800 Independence Avenue, SW., Washington, DC 20591; telephone: (202) 267–9255.

SUPPLEMENTARY INFORMATION:

Comments Invited

Interested parties are invited to participate in this proposed rulemaking by submitting such written data, views, or arguments as they may desire. Comments that provide the factual basis supporting the views and suggestions presented are particularly helpful in developing reasoned regulatory decisions on the proposal. Comments are specifically invited on the overall regulatory, aeronautical, economic, environmental, and energy-related aspects of the proposal. Communications should identify the airspace docket number and be submitted in triplicate to the address listed above. Commenters wishing the FAA to acknowledge receipt of their comments on this notice must submit with those comments a self-addressed, stamped postcard on which the following statement is made: "Comments to Airspace Docket No. 93–AGL–23." The postcard will be date/time stamped and returned to the commenter. All communications received on or before the specified closing data for comments will be considered before taking action on the proposed rule. The proposal contained in this notice may be changed in light of comments received. All comments submitted will be available for examination in the Rules Docket both before and after the closing date for comments. A report summarizing each substantive public contact with FAA personnel concerned with this rulemaking will be filed in the docket.

Availability of NPRM's

Any person may obtain a copy of this Notice of Proposed Rulemaking (NPRM) by submitting a request to the Federal Aviation Administration, Office of Public Affairs, Attention: Public Inquiry Center, APA–220, 800 Independence Avenue, SW., Washington, DC 20591, or by calling (202) 267–3485. Communications must identify the notice number of this NPRM. Persons interested in being placed on a mailing list for future NPRM's should also request a copy of Advisory Circular No. 11–2A, which describes the application procedure.

The Proposal

The FAA is considering an amendment to part 71 of the Federal Aviation Regulations (14 CFR part 71) to alter VOR Federal Airways V–172 and V–177 located in Wisconsin and Illinois. To expedite the flow of traffic, direct routings are desirable for aircraft arriving at satellite airports in the Chicago metropolitan area. V–172 would be realigned from the newly relocated ELGIN, IL, intersection direct to the Du Page, IL, (DPA) Very High Frequency Omnidirectional Range/Distance Measuring Equipment facility. V–177, heading southeast from the Janesville, WI, (JVL) Very High Frequency Omnidirectional Range/Tactical Air Navigation (VORTAC) facility toward the Joliet, IL, (JOT) VORTAC currently doglegs over the Rockford, IL, (RFD) VORTAC. Realigning V–177 would provide a direct route between the Janesville and Joliet VORTACs and eliminate the dogleg over the Rockford VORTAC. Domestic VOR Federal airways are published in paragraph 6010(a) of FAA Order 7400.9A dated June 17, 1993, and effective September 16, 1993, which is incorporated by reference in 14 CFR 71.1 (58 FR 36298; July 6, 1993). The airways listed in this document would be published subsequently in the Order.

The FAA has determined that this proposed 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, when promulgated, 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 71

Airspace, Incorporation by reference, Navigation (air).

The Proposed Amendment

In consideration of the foregoing, the Federal Aviation Administration proposes to amend 14 CFR part 71 as follows:

PART 71—[AMENDED]

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

Authority: 49 U.S.C. app. 1348(a), 1354(a), 1510; E.O. 10854, 24 FR 9565, 3 CFR, 1959–1963 Comp., p. 389; 49 U.S.C. 106(g); 14 CFR 11.69.

§ 71.1 [Amended]

2. The incorporation by reference in 14 CFR 71.1 of the Federal Aviation Administration Order 7400.9A, Airspace Designations and Reporting Points, dated June 17, 1993, and effective September 16, 1993, is amended as follows:

Paragraph 6010(a)—Domestic VOR Federal Airways

* * * * *

V–172 [Revised]

From North Platte, NE, via INT North Platte 073° and Wolbach, NE, 266° radials; Wolbach; Columbus, NE; Omaha, NE, INT Omaha 066° and Newton, IA, 262° radials; Newton; Cedar Rapids, IA; Polo, IL; INT Polo 088°T(085°M) and Du Page, IL, 293°T(291°M) radials; Du Page.

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V–177 [Revised]

From Joliet, IL, via Janesville, WI; Madison, WI; Stevens Point, WI; Wausau, WI; Hayward, WI; Duluth, MN; to Ely, MN.

* * * * *

Issued in Washington, DC, on April 7, 1994.

Harold W. Becker,

Manager, Airspace-Rules and Aeronautical Information Division.

[FR Doc. 94–9427 Filed 4–18–94; 8:45 am]

BILLING CODE 4910–13–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 343

[Docket No. 77N–094U]

RIN 0905–AA06

Internal Analgesic, Antipyretic, and Antirheumatic Drug Products for Over-the-Counter Human Use; Proposed Amendment to Tentative Final Monograph; Reopening of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of proposed rulemaking; reopening of comment period.

SUMMARY: The Food and Drug Administration (FDA) is reopening to May 4, 1994, the comment period on the notice of proposed rulemaking pertaining to combinations of over-the-counter (OTC) internal analgesic and antacid ingredients, specifically sodium bicarbonate used as an antacid active ingredient (February 2, 1994, 59 FR 5068). FDA is taking this action in response to a request to extend the comment period for an additional 30 days to allow more time to comment on this proposal. The comment period for these issues closed on April 4, 1994. This proposal is part of the ongoing review of OTC drug products conducted by FDA.

DATES: Written comments by May 4, 1994.

ADDRESSES: Written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, rm. 1–23, 12420 Parklawn Dr., Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: William E. Gilbertson, Center for Drug Evaluation and Research (HFD–810), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–594–5000.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of February 2, 1994 (59 FR 5068), FDA issued a notice of proposed rulemaking to amend the tentative final monograph for OTC internal analgesic, antipyretic, and antirheumatic drug products (hereinafter referred to as the February 1994, proposal). This proposal affected combinations of internal analgesic and antacid ingredients, specifically sodium bicarbonate used as an antacid active ingredient. As proposed, combination drug products intended to be dissolved in liquid prior to administration, such as powders and effervescent granules or tablets, would not be allowed to make a claim for "relief of overindulgence in food and drink" or a claim for "relief of hangover." FDA issued the February 1994, proposal after receiving reports of gastric (stomach) rupture following ingestion of sodium bicarbonate to relieve gastrointestinal distress. Interested persons were given until April 4, 1994, to submit comments on the proposal.

On March 28, 1994, the Nonprescription Drug Manufacturers Association (NDMA), a trade association representing the manufacturers and distributors of OTC medicines, requested that the comment period be extended for at least 30 days. NDMA stated that the extension is necessary to provide sufficient time to develop comprehensive comments to submit to the agency. NDMA explained that most companies that distribute OTC drug products containing sodium bicarbonate as an active ingredient were not anticipating changes in the internal analgesic, antipyretic, and antirheumatic tentative final monograph and the February 1994, proposal provided the first opportunity for many affected parties to become aware of the agency's concerns and intentions for amending the tentative final monograph. NDMA stated that interested parties have had insufficient time to acquire the reference materials cited by FDA, to retrieve relevant product experience reports, and to research thoroughly the issues raised in FDA's February 1994, proposal. NDMA also mentioned that the time period for the comments includes several days on which many company employees will not be working because of religious holidays.

FDA has carefully considered the request and believes that this additional time for comment is in the public interest. Accordingly, the comment period is reopened to May 4, 1994.

Interested persons may, on or before May 4, 1994, submit to the Dockets Management Branch (address above) written comments regarding OTC internal analgesic, antipyretic, and antirheumatic drug products containing sodium bicarbonate as an antacid active ingredient. Three copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the office above between 9 a.m. and 4 p.m., Monday through Friday.

Dated: April 13, 1994.

Michael R. Taylor,

Deputy Commissioner for Policy.

[FR Doc. 94–9354 Filed 4–18–94; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF LABOR

Occupational Safety and Health Administration

29 CFR Part 1903

[Docket No. C–03]

Abatement Verification

AGENCY: Occupational Safety and Health Administration (OSHA), Labor.

ACTION: Notice of proposed rulemaking.

SUMMARY: OSHA is developing a regulation requiring employers to certify abatement and submit abatement plans and progress reports as a result of OSHA citations. In addition, OSHA is proposing the placement of a tag on cited equipment to alert affected employees that a hazardous condition exists while abatement is being accomplished. Violation of the regulation would result in civil penalties as prescribed by section 17 of the Occupational Safety and Health Act of 1970. This notice invites interested parties to submit comments and recommendations on the issues detailed in this document, as well as other pertinent issues. All the information received in response to this notice will be carefully reviewed. The comments received will assist OSHA in developing the final regulation.

DATES: Written comments on the notice of proposed rulemaking must be postmarked no later than July 18, 1994.

ADDRESSES: Comments and information should be submitted in quadruplicate to the Docket Officer, Docket No. C–03, Occupational Safety and Health Administration, room N–2625, U.S. Department of Labor, 200 Constitution Ave. NW., Washington, DC 20210; telephone: (202) 219–7894.

FOR FURTHER INFORMATION CONTACT: Mr. James Foster, Occupational Safety and Health Administration, Office of Public Affairs, room N–3647, U.S. Department of Labor, 200 Constitution Ave. NW., Washington, DC 20210; telephone: (202) 219–8151.

SUPPLEMENTARY INFORMATION: The purpose of this proposed rule is to require employers to inform OSHA and their employees about measures they will take or have taken in response to OSHA citations, as well as to inform employees about OSHA citations and the alleged safety or health hazards described therein.

I. Background

Under the Occupational Safety and Health Act of 1970 (i.e., "the Act" or "the OSH Act"), 29 U.S.C. 651 *et. seq.*, OSHA inspects workplaces to determine whether employers are complying with OSHA standards and other statutory or regulatory requirements. If OSHA believes that an employer has committed a violation, a citation is issued. The citation will reference the requirement allegedly violated, the alleged violation, and note the proposed penalty and a date by which the violation is to be corrected, i.e., the abatement date. Section 9(a), 29 U.S.C. 658(a).