Airspace Management, ATA–400, 800 Independence Avenue, SW., Washington, DC 20591, or by calling (202) 267–8783. Communications must identify both docket numbers for this notice. Persons interested in being placed on a mailing list for future NPRM's should contact the FAA's Office of Rulemaking, (202) 267–9677, to request a copy of Advisory Circular No. 11–2A, Notice of Proposed Rulemaking Distribution System, which describes the application procedure.

The Proposal

The FAA is considering an amendment to 14 CFR part 71 to modify Class E airspace at South Bend, IN, for South Bend Regional Airport. Controlled airspace extending upward from 700 feet or more above the surface of the earth is needed to contain aircraft executing instrument approach procedures. The area would be depicted on appropriate aeronautical charts. Class E airspace areas extending upward from 700 feet or more above the surface of the earth are published in paragraph 6005 of FAA Order 7400.9K dated August 30, 2002, and effective September 16, 2002, which is incorporated by reference in 14 CFR 71.1. The Class E designations listed in this document would be published subsequently in the Order.

The FAA has determined that this proposed regulation only involves an establishment body of technical regulations for which frequent and routine amendments are necessary to keep them operationally current. Therefore, this proposed 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 proposed rule 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

Accordingly, pursuant to the authority delegated to me, the Federal Aviation Administration proposes to amend 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 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 the Federal Aviation Administration Order 7400.9K, Airspace Designations and Reporting Points, dated August 30, 2002, and effective September 16, 2002, is amended as follows:

* * * * *

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

* * * * *

AGL IN E5 South Bend, IN [Revised]

South Bend, South Bend Regional Airport, IN
(Lat. 41°42′31″ N., long. 86°19′02″ W.)
Niles, Jerry Tyler Memorial Airport, MI
(Lat. 41°50′09″ N., long. 86°13′31″ W.)
Gipper VORTAC
(Lat. 41°46′07″ N., long. 86°19′06″ W.)
That airspace extending upward from 700 feet above the surface within an 8.0-mile radius of South Bend Regional Airport and within 4.4 miles south and 7 miles north of the South Bend ILS localizer east course, extending from South Bend Regional Airport to 10.5 miles east of the ILS outer marker and within 4.4 miles west and 7 miles east of the Gipper VORTAC 001° radial, extending from the South Bend Regional Airport to 10.5 miles north of the VOR and within a 6.4-mile radius of the Jerry Tyler Memorial Airport, excluding that airspace within the Dowagiac, MI, Class E airspace area.

* * * * *

Issued in Des Plaines, Illinois on April 1, 2003.

Nancy B. Shelton,

Manager, Air Traffic Division, Great Lakes Region.

[FR Doc. 03–9729 Filed 4–18–03; 8:45 am]

BILLING CODE 4910–13–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 111 and 112

RIN 0910–AB88

[Docket No. 96N–0417]

Dietary Supplements; Current Good Manufacturing Practice Proposed Regulation; Notice of Public Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; satellite downlink public meeting.

SUMMARY: The Food and Drug Administration (FDA) is announcing a public meeting (via satellite downlink) to discuss the proposed rule on current good manufacturing practice in manufacturing, packing, or holding dietary ingredients and dietary supplements that published in the **Federal Register** of March 13, 2003. This satellite downlink public meeting is intended to provide clarification of the proposed rule and to explain how to submit comments on the proposed rule. This meeting will provide stakeholders, including small business, an opportunity to ask questions about the proposed rule by telephone, e-mail, or FAX. Questions also may be submitted in advance of the satellite downlink public meeting until the day before the downlink (see **FOR FURTHER INFORMATION CONTACT** section of this document).

DATES: The public meeting via satellite downlink will be held on May 9, 2003, from 12:30 p.m. to 3:30 p.m. eastern daylight time.

ADDRESSES: Downlink sites for the public meeting will be Food and Drug Administration District and Regional Offices for FDA employees. Small businesses wishing to view the satellite downlink should contact their Regional Small Business Representative. Regional representatives are listed at the Office of Regulatory Affairs' Web site at: *http://www.fda.gov/ora/fed_state/Small_business/sb_guide/smbusrep.htm*, or go to *http://www.fda.gov/ora/fed_state/events/default.htm* for a list of public viewing sites.

State and local counterparts who wish to participate may consider any local viewing location that has access to a C-band steerable dish.

Viewers with access to a steerable dish capable of receiving a C-band satellite signal may wish to tune this meeting in themselves. Tuning coordinates and course materials will be

placed on the Center for Food Safety and Applied Nutrition (CFSAN) Web site at: *http://www.cfsan.fda.gov/~dms/supplmnt.html* when available.

FOR FURTHER INFORMATION CONTACT: Bradford W. Williams, Center for Food Safety and Applied Nutrition (HFS–810), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, telephone: 301–436–1440, FAX: 301–436–2636, e-mail: *Brad.Williams@cfsan.fda.gov* for general questions about the downlink and submission of advance questions.

SUPPLEMENTARY INFORMATION:

I. Background

The Dietary Supplement Health and Education Act of 1994 (DSHEA) (Public Law 103–417) amended the Federal Food, Drug, and Cosmetic Act to, among other things, provide FDA with express statutory authority to prescribe current good manufacturing practice (CGMP) regulations for dietary supplements (21 U.S.C. 342(g)). In the **Federal Register** of March 13, 2003 (68 FR 12157), FDA published a proposed rule entitled ''Current Good Manufacturing Practice in Manufacturing, Packing, or Holding Dietary Ingredients and Dietary Supplements'' to establish CGMPs that include provisions on manufacturing, packaging, labeling, testing, quality control, releasing for distribution, and holding of dietary ingredients and dietary supplements. The proposed CGMPs are intended to help ensure that manufacturing, packing, and holding practices will not result in an adulterated or misbranded dietary supplement.

This downlink meeting will provide an opportunity to brief stakeholders on the proposed rule and allow them to ask questions about the proposed rule. It is also intended to fulfill part of the outreach requirements of the Small Business Regulatory Enforcement Fairness Act of 1996. The half-day meeting will focus on information for manufacturers, both large and small, with an emphasis on assistance to small firms. Small firms are encouraged to view and participate in this downlink meeting.

II. Agenda

The agenda will include an overview of the proposed rule with the following specific topics: (1) Personnel, (2) physical plant, (3) equipment and utensils, (4) production and process controls, (5) holding and distribution, (6) consumer complaints, and (7) recordkeeping. In addition to explaining the content of the proposed rule, we will instruct participants on the process for submitting comments. We also will discuss the types of comments and supporting information that would be most helpful to the agency in developing a final rule. Lastly, the meeting will describe how the Small Business Administration (SBA) can help small firms that might be affected by the proposed rule.

The primary intended audience is dietary ingredient and dietary supplement manufacturers, packagers, distributors, and holders, including small businesses, their representatives and consultants; Federal, State and local representatives; and FDA small business representatives and other interested FDA staff. Viewers are encouraged to watch the satellite program and participate in the question and answer periods. Any interested parties with access to a satellite dish may view the downlink directly. For specific technical details, including tuning coordinates, check the CFSAN Web site at: *http://www.cfsan.fda.gov/~dms/supplmnt.html* under ''Recent Announcements'' before the meeting.

Before the broadcast, we suggest that interested parties read the section in the March 13, 2003 (68 FR 12157), proposed rule entitled ''Proposal Highlights and Request for Comments,'' as well as the background document, fact sheet and the guidance for small businesses that are located at the CFSAN Web site noted above. In addition, a promotional flyer and specific technical tuning instructions will be added to the CFSAN Web site in the near future.

Questions may be submitted in advance of the satellite downlink public meeting until the day before the meeting (see the **FOR FURTHER INFORMATION CONTACT** section of this document).

III. Transcripts

A transcript of the program and all questions/answers will be added to docket 96N–0417 and may be examined at 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, as well as on the CFSAN Web site. You may request a transcript of the public meeting from the Freedom of Information Office (HFI–35), Food and Drug Administration, 5600 Fishers Lane, rm. 12A–16, Rockville, MD 20857, approximately 3 weeks after the meeting at a cost of 10 cents per page. In addition, a videotape of the satellite downlink public meeting will be available for viewing after the broadcast at the FDA Dockets Management Branch.

IV. Comments

To submit written comments on the proposed rule that published in the **Federal Register** of March 13, 2003, please follow the instructions in the ''Request for Comment'' section of that document (68 FR 12157 at 12248), a copy of which may be found at CFSAN's Web site at: *http://www.cfsan.fda.gov/~dms/supplmnt.html*.

Dated: April 15, 2003.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 03–9660 Filed 4–18–03; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF LABOR

Occupational Safety and Health Administration

29 CFR Parts 1910, 1915, and 1926

[Docket No. S–778–A]

RIN 1218–AB 81

Standards Improvement Project—Phase II

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

ACTION: Proposed rule; notice of hearing.

SUMMARY: On October 31, 2002, OSHA published a proposed rule entitled ''Standards Improvement Project—Phase II''. OSHA will convene an informal public hearing to receive testimony and documentary evidence relevant to the issues raised in the proposal. This action is in response to the interested parties who have requested the convening of a hearing.

DATES: *Informal public hearing.* The Agency will hold an informal public hearing beginning in Washington, DC, on July 8 to July 9, 2003. The hearing will commence at 10 a.m. on the first day, and at 9 a.m. on the second day and subsequent days if they prove necessary; however, the exact daily schedule is at the discretion of the presiding administrative law judge.

Notice of intention to appear to provide testimony at the informal public hearing. Interested parties who intend to present testimony at the informal public hearing must notify OSHA of their intention to do so no later than June 5, 2003.

Hearing testimony and documentary evidence. Interested parties who will be requesting more than 10 minutes to present their testimony, or who will be submitting documentary evidence at the hearing, must provide the Agency with copies of their full testimony and all