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—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. 289.

§71.1 [Amended]

2. The incorporation by reference in 14 CFR 71.1 of Federal Aviation Administration Order 7400.9G, Airspace Designations and Reporting Points, dated September 1, 1999, and effective September 16, 1999, is amended as follows:

Paragraph 5000 Class D airspace.

ASO MS D Greenwood, MS [Revised]

Greenwood-Leflore Airport, MS (Lat. 33°29′44″ N, long. 90°05′03″ W)

That airspace extending upward from the surface, to and including 2,700 feet MSL within a 4.4-mile radius of the Greenwood-Leflore Airport. This Class D airspace area is effective during the specific dates and times established in advance by a Notice to Airmen. The effective dates and times will thereafter be continuously published in the Airport/Facility Directory.

Paragraph 6004 Class E Airspace Areas Designated as an Extension to a Class D Airspace Area.

ASO MS E4 Greenwood, MS [Revised]

Greenwood-Leflore Airport, MS (Lat. 33°29′44″ N, long. 90°05′03″ W) Greenwood VORTAC

(Lat. 33°27′50" N, long. 90°16′38" W)

That airspace extending upward from the surface within 1.4 miles each side of the Greenwood VORTAC 079° radial, extending from the 4.4-mile radius of Greenwood-Leflore Airport to 4 miles east of the VORTAC. This Class E airspace area is effective during the specific dates and times established in advance by a Notice to Airmen. The effective dates and times will thereafter be continuously published in the Airport/Facility Directory.

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

ASO MS E5 Greenwood, MS [Revised]

Greenwood-Leflore Airport, MS (Lat. 33°29'44" N, long. 90°05'03" W) Greenwood VORTAC

(Lat. 33°27′50" N, long. 90°16′38" W)

That airspace extending upward from 700 feet above the surface within a 6.9-mile radius of Greenwood-Leflore Airport and within 1.2 miles each side of the Greenwood VORTAC 079° radial, extending from the 6.9-mile radius to 2 miles east of the VORTAC.

Issued in College Park, Georgia, on March 31, 2000.

Nancy B. Shelton,

Acting Manager, Air Traffic Division, Southern Region.

[FR Doc. 00–9216 Filed 4–18–00; 8:45 am] BILLING CODE 4910–13–M

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 71

[Airspace Docket No. 00-AGL-10]

Proposed Establishment of Class E Airspace; Minneapolis, Crystal Airport, MN

AGENCY: Federal Aviation Administration (FAA), DOT.

ACTION: Notice of proposed rulemaking.

summary: This action proposes to establish Class E airspace at Minneapolis, Crystal Airport, MN. Crystal Airport is served by Federal Aviation Regulations Part 135 air carrier operations. Controlled airspace extending upward from the surface is needed to contain aircraft executing instrument flight procedures and provide a safer operating environment when the control tower is closed. The airport meets the minimum communications and weather observation and reporting requirements for controlled airspace extending

upward from the surface. This action proposes to create controlled airspace with a 3.8-mile radius for this airport.

DATES: Comments must be received on or before May 22, 2000.

ADDRESSES: Send comments on the proposal in triplicate to: Federal Aviation Administration, Office of the Regional Counsel, AGL-7, Rules Docket No. 00-AGL-10, 2300 East Devon Avenue, Des Plaines, Illinois 60018.

The official docket may be examined in the Office of the Regional Counsel, Federal Aviation Administration, 2300 East Devon Avenue, Des Plaines, Illinois. An informal docket may also be examined during normal business hours at the Air Traffic Division, Airspace Branch, Federal Aviation Administration, 2300 East Devon Avenue, Des Plaines, Illinois.

FOR FURTHER INFORMATION CONTACT:

Denis C. Burke, Air Traffic Division, Airspace Branch, AGL–520, Federal Aviation Administration, 2300 East Devon Avenue, Des Plaines, Illinois 60018, telephone (847) 294–7568.

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 action must submit with those comments a self-addressed, stamped postcard on which the following statement is made: "Comments to Airspace Docket No. 00-AGL-10." The postcard will be date/ time stamped and returned to the commenter. All communications received on or before the specified closing date for comments will be considered before taking action on the proposed rule. The proposal contained in this action may be changed in light of comments received. All comments submitted will be available for examination in the Rules Docket, FAA, Great Lakes Region, Office of the Regional Counsel, 2300 East Devon Avenue, Des Plaines, Illinois, 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–230, 800 Independence Avenue, S.W., Washington, DC 20591, or by calling (202) 267–3484. Communications must identify the docket 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 14 CFR part 71 to establish Class E airspace at Minneapolis, Crystal Airport, MN, to accommodate FAR Part 135 (14 CFR part 135) air carrier aircraft executing instrument flight rules procedure during periods when the control tower is closed. The area would be depicted on appropriate aeronautical charts. Class E airspace designations for airspace areas extending upward from the surface of the earth are published in paragraph 6002 of FAA Order 7400.9G dated September 11, 1999, and effective September 16, 1999, which is incorporated by reference in 14 CFR 71.1. The Class E airspace 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 7400.9G, Airspace Designations and Reporting Points, dated September 11, 1999, and effective September 16, 1999, is amended as follows:

Paragraph 6002 Class E airspace designated as a surface area.

AGL MN E2 Minneapolis, Crystal Airport, MN [New]

Crystal Airport, MN (Lat. 45°08'42"N., long 93°12'41"W.)

Within a 3.8-mile radius of the Minneapolis, Anoka County-Blaine Airport. This Class E airspace area is effective during the specific dates and times established in advance by Notice to Airmen. The effective date and time will thereafter be continuously published in the Airport/Facility Directory.

Issued in Des Plaines, Illinois on March 22,

Christopher R. Blum,

Manager, Air Traffic Division. [FR Doc. 00–9215 Filed 4–18–00; 8:45 am] BILLING CODE 4910–13–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 864, 866, 868, 870, 872, 874, 876, 878, 884, 886, and 888

[Docket No. 99N-0035]

Medical Devices; Reclassification of 38 Preamendments Class III Devices into Class II

AGENCY: Food and Drug Administration, HHS.

ACTION: Proposed rule; reopening of comment period.

SUMMARY: The Food and Drug Administration (FDA) is reopening for 90 days the comment period for the submission of comments regarding 6 of the 38 devices proposed for reclassification from class III into class II. The proposed rule was published in the **Federal Register** of March 15, 1999 (64 FR 12774). The agency is taking this action in part in response to a request for more time to submit comments to FDA regarding several of the guidance documents that were not made available when the March 15, 1999, proposed rule was published. Elsewhere in this issue of the Federal Register, FDA is announcing the availability of these guidance documents for comment.

DATES: Submit written comments on the proposed rule by July 18, 2000.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Joseph M. Sheehan, Center for Devices and Radiological Health (HFZ–215), Food and Drug Administration, 1350 Piccard Dr., Rockville, MD 20850, 301–827–2974.

SUPPLEMENTARY INFORMATION:

I. Background

In the **Federal Register** of March 15, 1999 (64 FR 12774), FDA published a proposed rule to reclassify 38 preamendments class III devices into class II and to establish special controls for these devices. Interested persons were given until June 14, 1999, to comment on the proposed rule.

A trade association requested that FDA reopen the comment period for the following six devices: (1) Vascular graft prosthesis of less than 6 millimeters diameter, (2) pacemaker lead adaptor, (3) annuloplasty ring, (4) cardiopulmonary bypass defoamer, (5) cardiopulmonary bypass arterial line blood filter, and (6) cardiopulmomonary bypass oxygenator. The request noted that FDA had not made the guidance documents that were proposed as special controls for these six devices available for comment through the agency's Good Guidance Practices (GGP's). The request further noted that it was impossible to comment on the proposed reclassification without the guidance documents being available. Therefore, the trade association requested that FDA extend the comment period until at least 90 days after the guidance documents are publicly available for comment.

FDA also identified an additional three devices for which the agency had