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 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).

Adoption of the Amendment

In consideration of the foregoing, the Federal Aviation Administration amends 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 7400.9E, Airspace Designations and Reporting Points, dated September 10, 1997, and effective September 16, 1997, is amended as follows:

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

* * * * *

AGL ND E5 Crosby, ND [New]

Crosby Municipal Airport, ND
(Lat. 48°55′45″ N., long. 103°17′56″ W.)

That airspace extending upward from 700 feet above the surface within a 6.4-mile radius of the Crosby Municipal Airport, excluding that airspace north of lat. 49° 00′ 00″N (Canada/United States Boundary).

* * * * *

Issued in Des Plaines, Illinois, on August 25, 1998.

David B. Johnson,

Acting Manager, Air Traffic Division.

[FR Doc. 98–24290 Filed 9–9–98; 8:45 am]

BILLING CODE 4910–13–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 101

[Docket Nos. 98N–0426, 98N–0428, 98N–0427, 98N–0423, 98N–0424, 98N–0419, 98N–0422, 98N–0421, and 98N–0420]

Food Labeling: Health Claims; Reopening of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Interim final rules; reopening of comment period.

SUMMARY: The Food and Drug Administration (FDA) is reopening to October 8, 1998, the comment period for the nine interim final rules that appeared in the **Federal Register** of June 22, 1998 ((63 FR 34084), (63 FR 34092), (63 FR 34097), (63 FR 34101), (63 FR 34104), (63 FR 34107), (63 FR 34110), (63 FR 34112), and (63 FR 34115)). The rules prohibit the use on food labels of claims that are not appropriately based on authoritative statements from scientific bodies or that otherwise do not meet the specifications of new legislation. Interested persons were given until September 8, 1998, to comment on the interim final rules. This action is being taken in response to requests to reopen the comment period.

DATES: Written comments by October 8, 1998.

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: Christine J. Lewis, Center for Food Safety and Applied Nutrition (HFS–451), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–205–4168.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of June 22, 1998 ((63 FR 34084), (63 FR 34092), (63 FR 34097), (63 FR 34101), (63 FR 34104), (63 FR 34107), (63 FR 34110), (63 FR 34112), and (63 FR 34115)), FDA issued nine interim final rules prohibiting the use on food labels of claims that are not appropriately based on authoritative statements from scientific bodies or that otherwise do not meet the specifications of new legislation.

Interested persons were given until September 8, 1998, to comment on the rules. FDA has received several requests for extending the comment period. After evaluating these requests, the agency has decided to reopen the comment period on the interim final rules until October 8, 1998.

To be considered, written comments regarding the interim final rules must be received by October 8, 1998, by the Dockets Management Branch (address above). Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with Docket Nos. 98N–0426, 98N–0428, 98N–0427, 98N–0423, 98N–0424, 98N–0419, 98N–0422, 98N–0421, and 98N–0420. Received comments may be seen in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

Dated: September 4, 1998

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 98–24359 Filed 9–4–98; 4:34 pm]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 884

[Docket No. 97N–0335]

Obstetric and Gynecologic Devices; Reclassification and Classification of Medical Devices Used for In Vitro Fertilization and Related Assisted Reproduction Procedures

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is announcing that it is reclassifying instrumentation intended for use in in vitro fertilization (IVF) and related assisted reproduction technology (ART) procedures, including but not limited to gamete intrafallopian transfer (GIFT), embryo transfer (ET), and intracytoplasmic sperm injection (ICSI), from class III (premarket approval) to class II (special controls). FDA is also reclassifying assisted reproduction microscopes and microscope accessories from class III to class I. This reclassification is on the Secretary of the Department of Health and Human Services' (the Secretary's) own initiative based on new information. Accordingly, the order is being codified in the Code of Federal Regulations. Upon the effective date, this **Federal Register** document may be cited in the absence of an existing predicate device which would be used to support substantial equivalence. Elsewhere in this issue of the **Federal Register**, FDA is announcing the