

§ 39.13 [Amended]

2. Section 39.13 is amended by adding the following new airworthiness directive:

Boeing: Docket 2002–NM–287–AD.

Applicability: All Model 767–400ER series airplanes, certificated in any category.

Compliance: Required as indicated, unless accomplished previously.

To prevent fatigue cracking in the aft lower lug run-out region of the deflection control track, which could result in the loss of the secondary load path for the outboard flap, resulting in loss of the outboard flap and consequent reduced controllability of the airplane in the event that the primary load path also fails, accomplish the following:

Initial Inspection

(a) Before the accumulation of 12,000 total flight cycles, or within 1,200 flight cycles after the effective date of this AD, whichever occurs later, perform a high frequency eddy current (HFEC) inspection for cracks in the aft lower lug of the deflection control track on the outboard flap, in accordance with the Accomplishment Instructions of Boeing Alert Service Bulletin 767–27A0183, dated May 9, 2002.

Repetitive Inspections

(b) If no crack is detected during any HFEC inspection required in paragraph (a) of this AD, repeat the inspection at intervals not to exceed 1,200 flight cycles.

Corrective Action

(c) If any crack is detected during any HFEC inspection required by paragraph (a) of this AD, before further flight, replace the deflection control track with a new track assembly, in accordance with the Accomplishment Instructions in Boeing Alert Service Bulletin 767–27A0183, dated May 9, 2002. Within 12,000 flight cycles following the replacement, perform the HFEC inspection specified in paragraph (a) of this AD, and repeat inspections as specified in paragraph (b) of this AD.

Alternative Methods of Compliance

(d) In accordance with 14 CFR 39.19, the Manager, Seattle Aircraft Certification Office, FAA, is authorized to approve alternative methods of compliance for this AD.

Issued in Renton, Washington, on September 25, 2003.

Ali Bahrami,

Acting Manager, Transport Airplane Directorate, Aircraft Certification Service.
[FR Doc. 03–24842 Filed 9–30–03; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES**Food and Drug Administration****21 CFR Part 1**

[Docket Nos. 2002N–0276 and 2002N–0278]

Regulations Implementing Title III of the Public Health Security and Bioterrorism Preparedness and Response Act of 2002; Notice of Public Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Announcement of satellite downlink public meeting.

SUMMARY: The Food and Drug Administration (FDA) is announcing a public meeting (via satellite downlink) to discuss final regulations implementing two sections in Title III of the Public Health Security and Bioterrorism Preparedness and Response Act of 2002 (Bioterrorism Act) regarding Registration of Food Facilities (Docket No. 2002N–0276) and Prior Notice of Imported Food Shipments (Docket No. 2002N–0278). FDA expects to publish shortly in the **Federal Register** final rules implementing each of these provisions. The purpose of the satellite downlink public meeting is to provide information on the rules to the public and to provide the public an opportunity to ask questions of clarification.

DATES: The satellite downlink public meeting will be held on Tuesday, October 28, 2003, from 1 p.m. to 3 p.m. eastern standard time. Questions submitted in advance must be received by the contact person by close of business (4:30 p.m.) on Friday, October 24, 2003.

ADDRESSES: See **SUPPLEMENTARY INFORMATION** for locations where the satellite downlink may be viewed. A written transcript of the meeting will be available for viewing at the Division of Dockets Management (DDM) (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, and through the Web site at <http://www.fda.gov/oc/bioterrorism/bioact.html>.

A copy of the videotaped meeting may also be viewed at DDM.

FOR FURTHER INFORMATION CONTACT:

Louis J. Carson, Center for Food Safety and Applied Nutrition (HFS–32), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 301–436–2277, FAX: 301–436–2605, e-mail: CFSAN-FSS@cfsan.fda.gov, for general questions about the downlink,

submission of advance questions, and requests for a videotaped version of the meeting. Registration for specific downlink locations should be directed to the appropriate contact person listed in table 1 in the **SUPPLEMENTARY INFORMATION** section of this document.

SUPPLEMENTARY INFORMATION:**I. Background**

The events of September 11, 2001, highlighted the need to enhance the security of the U.S. food supply. Congress responded by passing the Public Health Security and Bioterrorism Preparedness and Response Act of 2002 (Public Law 107–188), which was signed into law on June 12, 2002. The Bioterrorism Act includes four provisions in Title III (Protecting Safety and Security of Food and Drug Supply), Subtitle A (Protection of Food Supply) that require the Secretary of Health and Human Services, through FDA, to develop implementing regulations on an expedited basis. These four provisions are section 305 (Registration of Food Facilities); section 307 (Prior Notice of Imported Food Shipments); section 306 (Maintenance and Inspection of Records for Foods); and section 303 (Administrative Detention). FDA expects that the agency will soon publish in the **Federal Register** final rules to implement sections 305 and 307 of the Bioterrorism Act. During the satellite downlink public meeting, FDA will explain the final rules on registration of food facilities and prior notice of imported food shipments and will answer questions. The satellite downlink public meeting will be offered in English with French and Spanish translation, and will be simulcast live in English, French, and Spanish for Mexico and North, Central, and South America (including Hawaii and Alaska).

II. Submitting Questions

Interested persons may submit questions concerning the final rules in advance of the downlink meeting. The deadline for the submission of questions is provided in the **DATES** section of this document. Questions submitted in advance will be used by the session moderator to help clarify issues of concern and provide information about the final rules. The viewing audience may telephone or fax questions to the FDA participants during the live downlink.

FDA is planning a second satellite downlink meeting during which FDA will explain the final rules that FDA intends to publish later this year to implement sections 306 and 303 of the Bioterrorism Act. That meeting will be