

Executive Order 12866; (2) is not a "significant rule" under DOT Regulatory Policies and Procedures (44 FR 11034, February 26, 1979); and (3) will not have a significant economic impact, positive or negative, on a substantial number of small entities under the criteria of the Regulatory Flexibility Act. A final evaluation has been prepared for this action and it is contained in the Rules Docket. A copy of it may be obtained from the Rules Docket at the location provided under the caption **ADDRESSES**.

List of Subjects in 14 CFR Part 39

Air transportation, Aircraft, Aviation safety, Incorporation by reference, Safety.

Adoption of the Amendment

■ Accordingly, pursuant to the authority delegated to me by the Administrator, the Federal Aviation Administration amends part 39 of the Federal Aviation Regulations (14 CFR part 39) as follows:

PART 39—AIRWORTHINESS DIRECTIVES

■ 1. The authority citation for part 39 continues to read as follows:

Authority: 49 U.S.C. 106(g), 40113, 44701.

§ 39.13 [Amended]

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

2004-07-06 Boeing: Amendment 39-13550. Docket 2002-NM-335-AD.

Applicability: Model 707 and 720 series airplanes, as listed in Boeing 707 Alert Service Bulletin A3509, dated June 13, 2002; certificated in any category.

Compliance: Required as indicated, unless accomplished previously.

To detect and correct cracking and/or loss of the upper and lower barrel nuts and bolts that retain the aft trunnion support fitting, which could result in the collapse of the main landing gear upon landing, accomplish the following:

Service Bulletin References

(a) The term "service bulletin," as used in this AD, means the Accomplishment Instructions of Boeing 707 Alert Service Bulletin A3509, dated June 13, 2002.

Initial Inspection

(b) Within 60 days after the effective date of this AD, for each main landing gear, perform the inspection specified in paragraph (b)(1) of this AD and the torque check specified in paragraph (b)(2) of this AD, in accordance with the service bulletin.

(1) Perform a detailed inspection of the upper and lower barrel nuts and bolts that retain the aft trunnion support fitting for corrosion, cracks, and loose or missing nuts and bolts.

(2) Torque check the upper and lower bolts to verify the torque is within the range specified in Figure 2 of the service bulletin.

Repetitive Inspections

(c) If no corrosion, crack, or loose or missing nut or bolt is found, and the torque is found to be within the specified range, during the inspection and torque check specified in paragraph (b) of this AD, then repeat the actions specified in paragraph (b) of this AD thereafter at intervals not to exceed 60 days.

Corrective Actions

(d) If any corrosion, crack, or loose or missing nut or bolt is found, or if the torque is found not to be within the specified range, during the inspection and torque check specified in paragraph (b) of this AD: Before further flight, do the corrective actions specified in paragraphs (d)(1) through (d)(3) of this AD. Accomplishment of these actions constitutes terminating action for the repetitive inspections specified in paragraph (c) of this AD.

(1) Perform a detailed inspection of the aft trunnion bearing cap and aft trunnion support fitting for corrosion, in accordance with the service bulletin. If any corrosion is detected, before further flight, repair in accordance with the service bulletin.

(2) Perform a magnetic particle inspection of the aft trunnion bearing cap for cracks in accordance with Figure 3 of the service bulletin.

(i) If no crack is found, before further flight, reinstall the inspected aft trunnion bearing cap in accordance with the service bulletin.

(ii) If any crack is found, before further flight, replace the aft trunnion bearing cap with a new aft trunnion bearing cap in accordance with the service bulletin.

(3) Reinstall the main landing gear trunnion with new Inconel barrel nuts and bolts to retain the aft trunnion support fitting, in accordance with Figure 4 of the service bulletin.

Terminating Action

(e) Within one year after the effective date of this AD, for each main landing gear, replace the upper and lower steel barrel nuts and H-11 bolts that retain the aft trunnion support fitting with new Inconel barrel nuts and bolts as specified in paragraphs (d)(1) through (d)(3) of this AD. Accomplishment of these actions constitutes terminating action for the requirements of this AD.

Parts Installation

(f) As of the effective date of this AD, no person shall install a steel barrel nut with H-11 bolt to retain the aft trunnion support fitting, on any airplane.

Alternative Methods of Compliance

(g) 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.

Incorporation by Reference

(h) The actions shall be done in accordance with Boeing 707 Alert Service Bulletin A3509, dated June 13, 2002. This incorporation by reference was approved by

the Director of the Federal Register in accordance with 5 U.S.C. 552(a) and 1 CFR part 51. Copies may be obtained from Boeing Commercial Airplanes, P.O. Box 3707, Seattle, Washington 98124-2207. Copies may be inspected at the FAA, Transport Airplane Directorate, 1601 Lind Avenue, SW., Renton, Washington; or at the Office of the Federal Register, 800 North Capitol Street, NW., suite 700, Washington, DC.

Effective Date

(i) This amendment becomes effective on May 11, 2004.

Issued in Renton, Washington, on March 19, 2004.

Kevin M. Mullin,

Acting Manager, Transport Airplane Directorate, Aircraft Certification Service.

[FR Doc. 04-7126 Filed 4-5-04; 8:45 am]

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DEPARTMENT OF TRANSPORTATION

Bureau of Industry and Security

15 CFR Part 774

[Docket No. 031201299-3299-01]

RIN 0694-AC54

Removal of "National Security" controls from, and imposition of "Regional Stability" controls on, certain items on the Commerce Control List; Correction

AGENCY: Bureau of Industry and Security, Commerce.

ACTION: Final rule; correction.

SUMMARY: The Bureau of Industry and Security published in the **Federal Register** of March 30, 2004, a final rule that replaced national security export and reexport controls on certain items with regional stability controls. This document corrects two typographical errors that appeared in that rule.

DATES: This rule is effective March 30, 2004.

FOR FURTHER INFORMATION CONTACT: William Arvin, Regulatory Policy Division, Office of Exporter Services, Bureau of Export Administration, Telephone: (202) 482-0436.

SUPPLEMENTARY INFORMATION: The Bureau of Industry and Security published in the **Federal Register** of March 30, 2004 (69 FR 16478), a final rule that replaced national security export and reexport controls on certain items with regional stability controls. That document inadvertently misstated a cross reference to Export Control Classification Number 0A984 as 0984. It also misstated a reference to Country Chart column "AT Column 1" as "AT

Column". This document corrects those errors.

PART 774—[CORRECTED]

■ In rule FR Doc. 04–7005, published on March 30, 2004 (69 FR 16478), make the following corrections. On page 16480, the middle column, correct the note to Export Control Classification Number 0A018, paragraph .c to read as follows:

Note: 0A018.c does not control weapons used for hunting or sporting purposes that were not specifically designed for hunting or sporting purposes that were not specially designed for military use and are not of the fully automatic type, but see 0A984 concerning shotguns.

■ On page 16480, the third column, in the Reason for Control paragraph of the License Requirements section of Export Control Classification Number 0E918, correct the third line in the Country Chart column to read: AT Column 1.

Eileen Albanese,

Director, Office of Exporter Services.

[FR Doc. 04–7808 Filed 4–5–04; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 312

Emergency Use of an Investigational New Drug; Technical Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending its regulations to reflect a change in address for the agency contacts for submitting an investigational new drug application (IND) in an emergency situation. This action is editorial in nature and is intended to improve the accuracy of the agency's regulations.

DATES: This rule is effective April 6, 2004.

FOR FURTHER INFORMATION CONTACT: Mark I. Fow, Office of Emergency Operations (HFA–615), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–443–1240.

SUPPLEMENTARY INFORMATION: FDA is amending its regulations in part 312 (21 CFR part 312) to reflect a change in address for the agency contacts for submitting an IND in an emergency situation that does not allow time for submission of an IND in accordance

with § 312.23 or § 312.34. The current address for submission of investigational biological drugs in an emergency situation is the "Division of Biological Investigational New Drugs (HFB–230), Center for Biologics Evaluation and Research, 8800 Rockville Pike, Bethesda, MD 20892, 301–443–4864." The new address for investigational biological drugs regulated by the Center for Biologics Evaluation and Research is "Office of Communication, Training and Manufacturers Assistance (HFM–40), Center for Biologics Evaluation and Research, 301–827–2000." The current contact for submission of all other investigational drugs in an emergency situation is the "Document Management and Reporting Branch (HFD–53), Center for Drug Evaluation and Research, 5600 Fishers Lane, Rockville, MD 20857, 301–443–4320." The new contact is the "Division of Drug Information (HFD–240), Center for Drug Evaluation and Research, 301–827–4570." The current contact for submitting requests for the Center for Biologics Evaluation and Research or the Center for Drug Evaluation and Research regulated products after normal working hours, eastern standard time, in an emergency situation is "FDA Division of Emergency and Epidemiological Operations, 202–857–8400." The new contact is "FDA Office of Emergency Operations (HFA–615), 301–443–1240."

Publication of this document constitutes final action on these changes under the Administrative Procedure Act (5 U.S.C. 553). Notice and public procedure are unnecessary because FDA is merely correcting nonsubstantive errors.

This rule does not meet the definition of "rule" in 5 U.S.C. 804(3)(A) because it is a rule of "particular applicability." Therefore, it is not subject to the congressional review requirements in 5 U.S.C. 801–808.

List of Subjects in 21 CFR Part 312

Drugs, Exports, Imports, Investigations, Labeling, Medical research, Reporting and recordkeeping requirements, Safety.

■ Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR part 312 is amended as follows:

PART 312—INVESTIGATIONAL NEW DRUG APPLICATION

■ 1. The authority citation for 21 CFR part 312 continues to read as follows:

Authority: 21 U.S.C. 321, 331, 351, 352, 353, 355, 371; 42 U.S.C. 262.

■ 2. Section 312.36 is revised to read as follows:

§ 312.36 Emergency use of an investigational new drug (IND).

Need for an investigational drug may arise in an emergency situation that does not allow time for submission of an IND in accordance with § 312.23 or § 312.34. In such a case, FDA may authorize shipment of the drug for a specified use in advance of submission of an IND. A request for such authorization may be transmitted to FDA by telephone or other rapid communication means. For investigational biological drugs regulated by the Center for Biologics Evaluation and Research, the request should be directed to the Office of Communication, Training and Manufacturers Assistance (HFM–40), Center for Biologics Evaluation and Research, 301–827–2000. For all other investigational drugs, the request for authorization should be directed to the Division of Drug Information (HFD–240), Center for Drug Evaluation and Research, 301–827–4570. After normal working hours, eastern standard time, the request should be directed to the FDA Office of Emergency Operations (HFA–615), 301–443–1240. Except in extraordinary circumstances, such authorization will be conditioned on the sponsor making an appropriate IND submission as soon as practicable after receiving the authorization.

Dated: March 31, 2004.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 04–7734 Filed 4–5–04; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HOMELAND SECURITY

Coast Guard

33 CFR Parts 101 and 104

[USCG–2004–17350]

Interpretation of International Voyage for Security Regulations

AGENCY: Coast Guard, DHS.

ACTION: Notice of interpretation.

SUMMARY: The Coast Guard is issuing an interpretation of the term "international voyage" as it is used in our recently-issued maritime security regulations. This interpretation will assist U.S. flag vessels operating in the waters of a foreign country in determining whether they must comply with the new International Ship and Port Facility