the requirements specified in paragraphs (l)(10)(i) and (l)(10)(ii) of this AD.
(5) CF34–BJ S/B 73–0062, Revision 01, dated July 1, 2008, or earlier issue, meet the requirements specified in paragraphs (l)(11) and (l)(12) of this AD.

Installation Prohibitions

(n) After the effective date of this AD:
(1) Do not install any fan blade into any CF34–3A1 engine with fan drive shaft, P/N 6036T78P02, with an airworthiness limitation section fan drive shaft life limit of 22,000 CSN if that fan blade:
(i) Was installed in a CF34–3A1 engine with fan drive shaft, P/N 6036T78P02, with an airworthiness limitation section fan drive shaft life limit of 15,000 CSN if that fan blade:
(ii) Is listed in Appendix A of GEAE SB No. CF34–BJ S/B 72–0230, Revision 01, dated July 30, 2008; or
(2) Do not install any fan blade into any CF34–3A1 engine with fan drive shaft, P/N 6036T78P02, with an airworthiness limitation section fan drive shaft life limit of 15,000 CSN if that fan blade:

Table 1—Material Incorporated by Reference

<table>
<thead>
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<th>Service Bulletin No.</th>
<th>Page</th>
<th>Revision</th>
<th>Date</th>
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(1) The Director of the Federal Register previously approved the incorporation by reference of this service information under § 52.10(a) of 5 U.S.C. 552(a) and 1 CFR part 51, as of January 4, 2010.

(2) For service information identified in this AD, contact General Electric Company, GE–Aviation, Room 285, 1 Neumann Way, Cincinnati, OH 45215, telephone (513) 552–3272; fax (513) 552–3329; e-mail: geae.aoc@ge.com.

(3) You may review copies at the FAA, New England Region, 12 New England Executive Park, Burlington, MA; or at the National Archives and Records Administration (NARA). For information on the availability of this material at NARA, call (202) 741–6030, or go to: http://www.archives.gov/federal-register/cfr/ibr-locations.html.

Issued in Burlington, Massachusetts, on December 29, 2009.

Francis A. Favara,
Manager, Engine and Propeller Directorate,
Aircraft Certification Service.

[FR Doc. E9–31274 Filed 1–7–10; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF COMMERCE

Bureau of Industry and Security

15 CFR Part 738

[RIN 0694–AE62]

Amendments to the Export Administration Regulations (EAR) Based Upon the Accession of Albania and Croatia to Formal Membership in the North Atlantic Treaty Organization (NATO)

Correction

In rule document E9–30484 beginning on page 68142 in the issue of Wednesday, December 23, 2009, make the following correction:
## Supplement No. 1 to Part 738—Commerce Country Chart

### [Reason for control]

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<th>Countries</th>
<th>Nuclear non-proliferation</th>
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<th>Missile Tech</th>
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2 See §742.4(a) for special provisions that apply to exports and reexports to these countries of certain thermal imaging cameras.

3 See §742.6(a)(3) for special provisions that apply to military commodities that are subject to ECCN OA919.

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

**21 CFR Part 529**

[Docket No. FDA–2009–N–0665]

**Certain Other Dosage Form New Animal Drugs; Sevoflurane**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Final rule.

**SUMMARY:** The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect approval of an abbreviated new animal drug application (ANADA) filed by Halocarbon Products Corp. The ANADA provides for the use of sevoflurane inhalant anesthetic in dogs.

**DATES:** This rule is effective January 8, 2010.

**FOR FURTHER INFORMATION CONTACT:** John K. Harshman, Center for Veterinary Medicine (HFV–170), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 240–276–8197, e-mail: john.harshman@fda.hhs.gov.

**SUPPLEMENTARY INFORMATION:** Halocarbon Products Corp., 887 Kinderkamack Rd., River Edge, NJ 07661, filed ANADA 200–467 that provides for use of Sevoflurane, an inhalant anesthetic, in dogs. Halocarbon Products Corp.’s Sevoflurane is approved as a generic copy of SEVOFLO (sevoflurane), sponsored by Abbott Laboratories, under NADA 141–103. The ANADA is approved as of November 27, 2009, and the regulations are amended in §529.2150 to reflect the approval.

In accordance with the freedom of information provisions of 21 CFR part 20 and 21 CFR 514.11(e)(2)(ii), a summary of safety and effectiveness data and information submitted to support approval of this application may be seen in the Division of Dockets Management (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.

The agency has determined under 21 CFR 25.33 that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

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 529**

Animal drugs.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR part 529 is amended as follows:

**PART 529—CERTAIN OTHER DOSAGE FORM NEW ANIMAL DRUGS**

1. The authority citation for 21 CFR part 529 continues to read as follows:


§529.2150 [Amended]

2. In paragraph (b) of §529.2150, remove “Nos. 000074 and 060307” and in its place add “Nos. 000074, 012164, and 060307”.


Bernadette Dunham,
Director, Center for Veterinary Medicine.

**DEPARTMENT OF COMMERCE**

**National Oceanic and Atmospheric Administration**

**50 CFR Part 648**

[Docket No. 0907241164–91415–02]

**RIN 0648–AY09**

**Magnuson-Stevens Fishery Conservation and Management Act Provisions; Fisheries of the Northeastern United States**

**AGENCY:** National Marine Fisheries Service (NMFS), National Oceanic and Atmospheric Administration (NOAA), Commerce.