For further information contact: James A. Lewis, Office of Strategic Trade and Foreign Policy Controls, Bureau of Export Administration, Telephone: (202) 482-4196.

SUMMARY: The Bureau of Export Administration (BXA) is amending the Export Administration Regulations (EAR) by reinstating provisions of License Exception AVS for temporary reexports to Libya of foreign registered aircraft subject to the EAR. This limited action is taken in response to suspended United Nations sanctions.

DATES: This rule is effective April 5, 1999.

The Export Administration Act (EAA) expired on August 20, 1994, the President invoked the International Emergency Economic Powers Act and continued in effect the EAR, and to the extent permitted by law, the provisions of the EAA, as amended, in Executive Order 12924 of August 19, 1994, as extended by the President's notices of extension of the EAR, and to the extent permissible under Executive Order 12612. Although the Export Administration Act (EAA) expired on August 20, 1994, the President invoked the International Emergency Economic Powers Act and continued in effect the EAR, and to the extent permitted by law, the provisions of the EAA, as amended, in Executive Order 12924 of August 19, 1994, as extended by the President's notices of extension of the EAR, and to the extent permissible under Executive Order 12612.

Two of the three offices (Export Processing Zone Administration and the Science-Based Industrial Park Administration) are in special economic zones and are responsible for the activity in their respective zones.
requires that a notice of proposed rulemaking and an opportunity for public comment be given for this rule. Because a notice of proposed rule making and opportunities for public comment are not required to be given for this rule by 5 U.S.C. 553, or by any other law, the analytical requirements of the Regulatory Flexibility Act, 5 U.S.C. 601 et seq., are inapplicable. Therefore, this regulation is issued in final form. Although there is no formal comment period, public comments on this regulation are welcome on a continuing basis. Comments should be submitted to Frank J. Ruggiero, Office of Export Services, Bureau of Export Administration, Department of Commerce, P.O. Box 273, Washington, D.C. 20044.

List of Subjects in 15 CFR Parts 746

Embargoes, Exports, Foreign trade, Reporting and recordkeeping requirements.

Accordingly, Part 746 of the Export Administration Regulations (15 CFR Parts 730–774) is amended to read as follows:

1. The authority citation for 15 CFR Part 746 is revised to read as follows:


PART 746—AMENDED

2. Section 746.4 is amended by revising paragraph (b)(2)(ii)(G) to read as follows:

§ 746.4 Libya

(b) * * * * * * * * * *

(ii) * * * * * * * * * *

(G) Aircraft and vessels (AVS) for vessels only (see § 740.15(c)(1) of the EAR), and temporary reexports of foreign registered aircraft (see § 740.15(a)(4) of the EAR).

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Iain S. Baird,

Deputy Assistant Secretary for Export Administration.

[FR Doc. 99-23785 Filed 9-10-99; 8:45 am]

BILLING CODE 3510-33-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 5

Delegations of Authority and Organization; Technical Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending the regulations for delegations of authority to correct position titles for delegates in the Center for Drug Evaluation and Research (CDER). This action is necessary to ensure the continued accuracy of the regulations.

EFFECTIVE DATE: September 13, 1999.

FOR FURTHER INFORMATION CONTACT: Leanne Cusumano, Center for Drug Evaluation and Research (HFD-007), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, or Donna G. Page, Division of Management Programs (HFA-340), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–594–2041, or

YELLOW: 21 CFR Part 5

3. Section 5.31 is amended by revising paragraph (f) (3) to read as follows:

§ 5.31 Petitions under part 10.

(3) The Director and Deputy Director, Office of Generic Drugs, Office of Pharmaceutical Science, CDER, except for those drug products listed in § 314.440(b) of this chapter, are authorized to issue responses to citizen petitions submitted under § 10.30 of this chapter seeking a determination of the suitability of an abbreviated new drug application for a drug product.

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4. Section 5.93 is amended by revising paragraph (b) to read as follows:

§ 5.93 Submission of and effective approval dates for abbreviated new drug applications and certain new drug applications.

(b) The Director and Deputy Director, Office of Generic Drugs, Office of Pharmaceutical Science, CDER.

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William K. Hubbard,

Senior Associate Commissioner for Policy, Planning and Legislation.

[FR Doc. 99–23683 Filed 9–10–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

New Animal Drugs for Use in Animal Feeds; Nicarbazin and Bambermycins

AGENCY: Food and Drug Administration, HHS.