Holy See
Hungary
Iceland
India
Indonesia
Iran
Ireland
Italy
Japan
Jordan
Kenya
Korea (Republic of)
Kuwait
Laos (P.D.R.)
Latvia
Lesotho
Lithuania
Luxembourg
Macedonia
Malawi
Maldives
Mali
Malta
Mauritius
Mauritania
Mexico
Micronesia
Moldova (Republic of)
Monaco
Mongolia
Morocco
Namibia
Nepal
Netherlands
New Zealand
Niger
Nigeria
Norway
Oman
Pakistan
Panama
Papua New Guinea
Paraguay
Peru
Philippines
Poland
Portugal
Qatar
Romania
Russian Federation
Saint Lucia
Saudi Arabia
Senegal
Seychelles
Singapore
Slovak Republic
Slovenia
South Africa
Spain
Sri Lanka
Sudan
Suriname
Swaziland
Sweden
Switzerland
Tajikistan
Tanzania
Togo
Trinidad and Tobago
Tunisia
Turkey
Turkmenistan
United Kingdom
Ukraine
United States
Uruguay
Uzbekistan
Venezuela
Vietnam
Zimbabwe

6. Supplement No. 3 to part 745 is amended by revising the title to the supplement, and the entry for Taiwan, to read as follows:

Supplement No. 3 to Part 745—Foreign Authorized Agencies Responsible for Issuing End-Use Certificates Pursuant to § 745.2

* * * * *

Taiwan [1]

Board of Foreign Trade, Ministry of Economic Affairs, 1 Hukou St., Taipei, Tel: (02) 2351–0271, Fax: (02) 2351–3603

Export Processing Zone Administration, Ministry of Economic Affairs, 600 Chiachang Rd., Nantze, Kaohsiung, Tel: (07) 361–1212, Fax: (07) 361–4348

Science-Based Industrial Park Administration, National Science Council, Executive Yuan, 2 Hsin-an Rd., Hsinchu, Tel: (03) 577–3311, Fax: (03) 577–6222

Dated: September 1, 1999.

R. Roger Majak,

Assistant Secretary for Export Administration.

[FR Doc. 99–23309 Filed 9–10–99; 8:45 am]

BILLING CODE 3510–33–P

DEPARTMENT OF COMMERCE

Bureau of Export Administration

15 CFR Part 746

[Docket No. 990827238–9238–01]

RIN 0694–AB94

Reexports to Libya of Foreign Registered Aircraft Subject to the Export Administration Regulations

AGENCY: Bureau of Export Administration, Commerce.

ACTION: Final rule.

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.

FOR FURTHER INFORMATION CONTACT: James A. Lewis, Office of Strategic Trade and Foreign Policy Controls, Bureau of Export Administration, Telephone: (202) 482–4196.

SUPPLEMENTARY INFORMATION:

Background

On April 5, 1999, the United Nations Security Council (UNSC) suspended the sanctions against Libya set forth in UNSC resolutions 748 and 883. In light of this suspension, the United States has taken action that will allow, under License Exception AVS, the temporary reexport to Libya of foreign registered aircraft subject to the EAR. Foreign registered aircraft meeting all the temporary sojourn requirements of License Exception AVS may fly from foreign countries to Libya without obtaining prior written authorization from BXA. This action is limited in scope and in no way impacts other U.S. sanctions against Libya. Note that License Exception AVS remains unavailable for U.S. registered aircraft.

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 August 15, 1995 (60 FR 42767), August 14, 1996 (61 FR 42527) August 13, 1997 (62 FR 43629), August 13, 1998 (63 FR 44121), and August 10, 1999 (64 FR 44101).

Rule Making Requirements

1. This final rule has been determined to be non-significant for purposes of E.O. 12866.

2. Notwithstanding any other provision of law, no person is required to respond to nor be subject to a penalty for failure to comply with a collection of information subject to the requirements of the Paperwork Reduction Act unless that collection of information displays a current valid OMB Control Number. This regulation does not involve any paperwork collections.

3. This rule does not contain policies with Federalism implications sufficient to warrant preparation of a Federalism assessment under Executive Order 12612.

4. The provisions of the Administrative Procedure Act requiring notice of proposed rulemaking, the opportunity for public participation, and a delay in effective date, are inapplicable because this regulation involves a military or foreign affairs function of the United States (see 5 U.S.C. 553(a)(1)). Further, no other law

[1] 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 Exporter 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:

Authority: 50 U.S.C. app. 2401 *et seq.*; 50 U.S.C. 1701 *et seq.*; 22 U.S.C. 287c; 22 U.S.C. 6004; E.O. 12854, 58 FR 36587, 3 CFR 1993 Comp., p. 614; E.O. 12918, 59 FR 28205, 3 CFR, 1994 Comp., p. 899; E.O. 12924, 59 FR 43437, 3 CFR, 1994 Comp., p.917; E.O. 13088, 63 FR 32109, 3 CFR, 1998 Comp., p. 191; E.O. 13121 of April 30, 1999, 64 FR 24021 (May 5, 1999); Notice of August 10, 1999, 64 FR 44101 (August 13, 1999).

PART 746—[AMENDED]

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

§ 746.4 Libya

* * * * *

(b) * * *

(2) * * *

(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).

* * * * *

Dated: September 7, 1999.

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, 301–594–2041, or
Donna G. Page, Division of Management Programs (HFA–340), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4816.

SUPPLEMENTARY INFORMATION: FDA is correcting its regulations in subpart B of part 5 (21 CFR part 5) in two sections that reflect incorrect position titles for delegates within CDER. In the **Federal Register** of January 17, 1997 (62 FR 2554), FDA amended the regulations for delegations of authority to update titles of CDER delegates and organizational components to reflect organizational restructuring. In two instances, the position titles for the Director and Deputy Director, Office of Generic Drugs (OGD), Office of Pharmaceutical Science (OPS), CDER were inadvertently changed to reflect the Director and Deputy Director, Division of Bioequivalence, OGD, OPS, CDER. Previously, the Director and Deputy Director, OGD, OPS, CDER held those authorities. The Director and Deputy Division Director of Bioequivalence titles should be removed.

List of Subjects in 21 CFR Part 5

Authority delegations (Government agencies), Imports, Organization and functions (Government agencies).

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR part 5 is amended as follows:

PART 5—DELEGATIONS OF AUTHORITY AND ORGANIZATION

1. The authority citation for 21 CFR part 5 continues to read as follows:

Authority: 5 U.S.C. 504, 552, App. 2; 7 U.S.C. 138a, 2271; 15 U.S.C. 638, 1261–1282, 3701–3711a; 15 U.S.C. 1451–1461; 21 U.S.C. 41–50, 61–63, 141–149, 321–394, 467f, 679(b), 801–886, 1031–1309; 35 U.S.C. 156; 42 U.S.C. 241, 242, 242a, 242l, 242n 243, 262, 263, 264, 265, 300u–300u–5, 300aa–1; 1395y, 3246b, 4332, 4831(a), 10007–10008; E.O. 11921, 41 FR 24294, 3 CFR, 1977 Comp., p. 124–131; E.O. 12591, 52 FR 13414, 3 CFR, 1988 Comp., p. 220–223.

§ 5.22 [Amended]

2. Section 5.22 *Certification of true copies and use of Departmental seal* is amended by removing paragraph (a) (13) (viii).

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

§ 5.31 Petitions under part 10.

(f) * * *

(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.

* * * * *

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.

* * * * *

Dated: September 7, 1999.

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.