of the Federal Aviation Regulations (14 CFR 21.197 and 21.199) to operate the airplane to a location where the requirements of this AD can be accomplished.

(f) An alternative method of compliance or adjustment of the initial or repetitive compliance times that provides an equivalent level of safety may be approved by the Manager, Airplane Certification Office (ACO), FAA, 2601 Meacham Boulevard, Fort Worth, Texas 76193–0150. The request shall be forwarded through an appropriate FAA Maintenance Inspector, who may add comments and then send it to the Manager, Fort Worth ACO. Alternative methods of compliance approved in accordance with AD 93–19–06 (superseded by this action) are not considered approved as alternative methods of compliance with this AD.

Note 4: Information concerning the existence of approved alternative methods of compliance with this AD, if any, may be obtained from the Fort Worth ACO.

(g) The inspections required by this AD shall be done in accordance with Fairchild Service Bulletin 26–56–20–042, Issued: November 28, 1988; Revised: February 7, 1991, Fairchild Service Bulletin 226–56–001, Issued: February 2, 1983; Revised: November 26, 1991, Fairchild Service Bulletin 227–56–001, Issued: February 2, 1983; Revised: November 26, 1991, Fairchild Service Bulletin 226–56–002, Issued: March 3, 1983; Revised: May 29, 1992, Fairchild Service Bulletin 227–56–002, Issued: January 5, 1984; Revised: May 29, 1992, and April 1, 1993, Fairchild Service Bulletin 226–56–003, Issued: September 13, 1984; Revised: November 2, 1989, Fairchild Service Bulletin 227–56–003, Issued: September 13, 1984; Revised: November 2, 1989, and Fairchild Service Bulletin 26–56–10–038, Issued: October 8, 1984; Revised: February 7, 1991, as applicable. This incorporation by reference was previously 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 Fairchild Aircraft, P.O. Box 790490, San Antonio, Texas 78279–0490. Copies may be inspected at the FAA, Central Region, Office of the Assistant Chief Counsel, Room 1558, 601 E. 12th Street, Kansas City, Missouri, or at the Office of the Federal Register, 800 North Capitol Street, NW., suite 700, Washington, DC.

(h) This amendment (39–9774) supersedes AD 93–19–06, Amendment 39–8705.

(i) This amendment (39–9774) becomes effective on November 14, 1996.

Issued in Kansas City, Missouri, on September 19, 1996.

Michael Gallagher,

Manager, Small Airplane Directorate, Aircraft Certification Service.

[FR Doc. 96–24886 Filed 9–27–96; 8:45 am]

BILLING CODE 4910–13–U

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

New Animal Drugs For Use In Animal Feeds; Monensin

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 a supplemental new animal drug application (NADA) filed by Elanco Animal Health, Division of Eli Lilly and Co. The supplemental NADA provides for monensin Type A medicated articles to be used to make free-choice Type C medicated feeds for pasture cattle weighing less than 400 pounds for increased rate of weight gain.

EFFECTIVE DATE: September 30, 1996.

FOR FURTHER INFORMATION CONTACT: Jack Caldwell, Center for Veterinary Medicine (HFV–126), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1638.

SUPPLEMENTARY INFORMATION: Elanco Animal Health, Division of Eli Lilly and Co., Lilly Corporate Center, Indianapolis, IN 46285, filed supplemental NADA 95–735, which provides for use of a monensin Type A medicated article to make a monensin Type C medicated feed/free-choice mineral granule containing 1620 grams monensin per ton (g/t) to be fed at 50 to 200 milligrams per head per day free-choice to pasture cattle (slaughter, stocker, feeder, and dairy and beef replacement heifers) for increased rate of weight gain.

The supplemental NADA provides for removal of the restriction concerning use of the product for animals weighing less than 400 pounds body weight. The supplemental NADA is approved as of September 30, 1996, and the regulations are amended in 21 CFR 558.355(f)(3)(x)(c) to reflect the approval.

Under section 512(c)(2)(F)(iii) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b(c)(2)(F)(iii)), approval of this supplemental NADA does not qualify for marketing exclusivity because the supplement does not contain reports of new clinical or field investigations (other than bioequivalence or residue studies) or new human food safety studies (other than bioequivalence or residue studies) essential to the approval and conducted or sponsored by the applicant.

Approval of this supplemental NADA does not require a freedom of information (FOI) summary because the approval relies on data and information filed to support a previously approved supplement. FOI summaries for prior approvals may be seen in the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857, from 9 a.m. to 4 p.m., Monday through Friday.

The agency has determined under 21 CFR 25.24(d)(1)(i) 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.

List of Subjects in 21 CFR Part 558

Animal drugs, Animal feeds

Therefore, under the Federal Food, Drug, and Cosmetic Act and under the authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR part 558 is amended as follows:

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

1. The authority citation for 21 CFR part 558 continues to read as follows:

Authority: Secs. 512, 701 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b, 371).

§ 558.355 [Amended]

2. Section 558.355 *Monensin* is amended in paragraph (f)(3)(x)(*c*) in the first sentence by removing the phrase ''weighing more than 400 pounds''.

Dated: September 3, 1996.

Robert C. Livingston,

Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.

[FR Doc. 96–24965 Filed 9–27–96; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

21 CFR Part 1313

[DEA Number 110F]

RIN 1117–AA21

Distribution of Chemical Import/Export Declaration

AGENCY: Drug Enforcement Administration (DEA), Justice.

ACTION: Final rule.

SUMMARY: DEA is amending its regulations to clarify the distribution requirements for the Precursor and Essential Chemical Import/Export Declaration (DEA Form 486). The regulations do not specify that a copy of the form must be provided to the United States Customs Service (Customs) on or before the day of exportation, as required in the instructions on the form. This amendment to the regulations will eliminate any possibility for confusion as to when the form must be provided to Customs.

EFFECTIVE DATE: November 29, 1996.

FOR FURTHER INFORMATION CONTACT: G. Thomas Gitchel, Chief, Liaison and Policy Section, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Telephone (202) 307–7297.

SUPPLEMENTARY INFORMATION: On September 23, 1993, DEA published a notice of proposed rulemaking (NPRM) in the Federal Register (58 FR 49455) regarding the distribution of the Precursor and Essential Chemical Import/Export Declaration (DEA Form 486). The NPRM pointed out a discrepancy between the instructions contained in the DEA Form 486 and the requirements of Title 21, Code of Federal Regulations (CFR), § 1313.23(c). The instructions require that Copy 3 of the form be provided to Customs *on or before the day of exportation.* Section 1313.23(c) provides the same instructions but omits the phrase *on or before the day of exportation.* To avoid the possibility of confusion regarding when Copy 3 of the form should be provided to Customs, DEA proposed to amend § 1313.23(c) to be consistent with the instructions contained in the form. In addition, while it could not be included as a requirement, DEA also proposed to include a suggestion in the section that the exporter submit the Shipper's Export Document on or before the day of exportation, in order to facilitate the uninterrupted export of the goods.

No comments or objections were received regarding the proposed amendment to the regulations. Therefore, DEA is amending 21 CFR 1313.23(c) to include the appropriate language to be consistent with the instructions contained in the DEA Form 486.

The Deputy Assistant Administrator, Office of Diversion Control, hereby certifies that this action will have no significant impact upon entities whose interests must be considered under the Regulatory Flexibility Act (5 U.S.C. 601 et seq.). This action does not impose any new requirements or burden on the regulated industry. This action is being taken to clarify the requirements regarding the distribution of the DEA Form 486.

This rule has been drafted and reviewed in accordance with Executive Order 12866. DEA has determined that this is not a significant regulatory action under the provisions of Executive Order 12866, section 3(f). This rule clarifies existing requirements and will prevent confusion that might cause delays in the export of chemicals.

This action has been analyzed in accordance with the principles and criteria in Executive Order 12612, and it has been determined that the rule does not have sufficient federalism implications to warrant the preparation of a Federalism Assessment.

List of Subjects in 21 CFR Part 1313

Drug traffic control, Exports, Imports, Reporting requirements.

For reasons set out above, 21 CFR part 1313 is amended as follows:

PART 1313—[AMENDED]

1. The authority citation for part 1313 continues to read as follows:

Authority: 21 U.S.C. 802, 830, 871(b), 971

2. Section 1313.23 is amended by revising paragraph (c) to read as follows:

§ 1313.23 Distribution of export declaration.

* * * * *

(c) Copy 3 shall be presented to the U.S. Customs Service at the port of exit for each export of a listed chemical or chemicals on or before the day of exportation, and when possible, along with the Shippers Export Declaration.

Dated: September 6, 1996.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control.

[FR Doc. 96–24944 Filed 9–27–96; 8:45 am]

BILLING CODE 4410–09–M

UNITED STATES INFORMATION AGENCY

22 CFR Part 505

Privacy Act Regulation

AGENCY: United States Information Agency.

ACTION: Interim final rule with request for comments.

SUMMARY: This interim final rule will amend the existing Privacy Act Regulation implementing the Privacy Act of 1974, as amended. Changes in the regulation are to reconcile them with changes in the law and Agency policy. The Agency expects the amended regulation will enable Privacy Act requesters to better understand how to make requests and how the Agency responds to such requests.

DATES EFFECTIVE: October 15, 1996. Comments regarding this interim final rule will be accepted until October 30, 1996.

ADDRESSES: Comments may be mailed to the FOIA/Privacy Act Officer, U.S. Information Agency, Room M–29, 301 4th Street, SW., Washington, DC. 20547.

FOR FURTHER INFORMATION CONTACT: FOIA/PA Unit, U.S. Information Agency, Room M–29, 301 4th Street, SW., Washington, DC. 20547; telephone (202) 619–5499.

SUPPLEMENTARY INFORMATION: The privacy of each individual is directly affected by the collection, maintenance, use and dissemination of personal information by Federal agencies. In order to protect the privacy of individuals identified in information systems maintained by Federal agencies, it is important to regulate the collection, maintenance, use and dissemination of such information by such agencies. Therefore it becomes paramount to ensure that regulations implementing the law are clear and readily understandable to the public.

List of Subjects in 22 CFR Part 505

Privacy.

For the reasons given in the preamble, Part 505 of Title 22 is revised to read as follows:

PART 505—PRIVACY ACT POLICIES AND PROCEDURES

Sec.

Authority: Pub. L. 93–579, 88 Stat. 1897; 5 U.S.C. 552a; 55 FR 31940, Aug. 6, 1990, as amended.

§ 505.1 Purpose and scope.

The United States Information Agency will protect individuals' privacy from