adopt the interim rule as a final rule without change.

The Regulatory Flexibility Act and Executive Order 12866

As discussed in the interim rule, since the amendments are not subject to the notice and public procedure requirements of the Administrative Procedure Act (5 U.S.C. 553), they are not subject to the Regulatory Flexibility Act (5 U.S.C. 601 *et seq.*). Also, because this document involves a foreign affairs function of the United States and implements an international agreement, it is not subject to the provisions of E.O. 12866.

Paperwork Reduction Act

The collections of information involved in this interim rule have already been approved by the Office of Management and Budget (OMB) in accordance with the Paperwork Reduction Act of 1995 (44 U.S.C. 3507) and assigned OMB Control Numbers 1515–0065 (Entry summary and continuation sheet) and 1515–0214 (General recordkeeping and record production requirements). This rule does not propose any substantive changes to the existing approved information collections.

An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless the collection of information displays a valid control number assigned by OMB.

List of Subjects

19 CFR Part 132

Agriculture and agricultural products, Customs duties and inspection, Quotas, Reporting and recordkeeping requirements.

19 CFR Part 163

Administrative practice and procedure, Customs duties and inspection, Imports, Reporting and recordkeeping requirements.

Amendments to the Regulations

Accordingly, the interim rule amending 19 CFR parts 132 and 163, which was published in the **Federal Register** at 65 FR 5430 on February 4, 2000, is adopted as a final rule without change.

Raymond W. Kelly,

Commissioner of Customs.

Approved: June 14, 2000.

John P. Simpson,

Deputy Assistant Secretary of the Treasury.

[FR Doc. 00–17927 Filed 7–13–00; 8:45 am]

BILLING CODE 4820–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 821, 895, and 900

[Docket No. 00N–1361]

Code of Federal Regulations; Technical Amendments

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendments.

SUMMARY: The Food and Drug Administration (FDA) is amending its regulations to correct some errors that have become incorporated into the regulations. This action is being taken to improve the accuracy of the regulations.

DATES: This rule is effective July 14, 2000.

FOR FURTHER INFORMATION CONTACT: LaJuana D. Caldwell, Office of Policy, Planning, and Legislation (HF–927), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7010.

SUPPLEMENTARY INFORMATION: FDA has discovered that errors have been incorporated into the agency's codified regulations for 21 CFR parts 821, 895, and 900. This document corrects those errors. Publication of this document constitutes final action under the Administrative Procedure Act (5 U.S.C. 553). FDA has determined that notice and public comment are unnecessary because this amendment is nonsubstantive.

List of Subjects

21 CFR Part 821

Imports, Medical devices, Reporting and recordkeeping requirements.

21 CFR Part 895

Administrative practice and procedure, Labeling, Medical devices.

21 CFR Part 900

Electronic products.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR parts 821, 895, and 900 are amended as follows:

PART 821—MEDICAL DEVICE TRACKING REQUIREMENTS

1. The authority citation for 21 CFR part 821 continues to read as follows:

Authority: 21 U.S.C. 331, 351, 352, 360, 360e, 360h, 360i, 371, 374.

§ 821.50 [Amended]

2. Section 821.50 *Availability* is amended in paragraph (a) by removing ''Form FD 482'' and by adding in its place ''Form FDA 482''.

PART 895—BANNED DEVICES

3. The authority citation for 21 CFR part 895 continues to read as follows:

Authority: 21 U.S.C. 352, 360f, 360h, 360i, 371.

§ 895.21 [Amended]

4. Section 895.21 *Procedures for banning a device* is amended in the fourth sentence of paragraph (d)(8) by removing ''201(y)'' and by adding in its place ''201(x)''.

PART 900—MAMMOGRAPHY

5. The authority citation for 21 CFR part 900 continues to read as follows:

Authority: 21 U.S.C. 360i, 360nn, 374(e); 42 U.S.C. 263b.

§ 900.12 [Amended]

6. Section 900.12 *Quality standards* is amended in paragraph (e)(5)(iii)(A)(*1*) by removing ''Cycles/millimeters'' and by adding in its place ''Cycles/millimeter'', and in the third sentence of paragraph (f)(3) by removing ''results and notifying'' and by adding in its place ''results and for notifying''.

Dated: June 27, 2000.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 00–17811 Filed 7–13–00; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

21 CFR Part 1308

[DEA–187F]

RIN 1117–AA51

Schedules of Controlled Substances: Exempt Anabolic Steroids Products

AGENCY: Drug Enforcement Administration, Department of Justice.

ACTION: Final rule.

SUMMARY: The Drug Enforcement Administration (DEA) published an interim rule with request for comments (65 FR 3124, Jan. 20, 2000, as corrected at 65 FR 5024, Feb. 2, 2000) which identified six anabolic steroid products as being exempt from certain regulatory provisions of the Controlled Substances Act (21 U.S.C. 801 *et seq.*) (CSA). No