DEPARTMENT OF THE TREASURY

Customs Service

19 CFR Parts 132 and 163 [T.D. 00-49]

RIN 1515-AC55

Export Certificates for Sugar-Containing Products Subject to Tariff-Rate Quota

AGENCY: Customs Service, Department of the Treasury.

ACTION: Final rule.

SUMMARY: This document adopts as a final rule, without change, the interim rule amending the Customs Regulations that was published in the Federal Register on February 4, 2000, as T.D. 00-7. The interim rule set forth the form and manner by which an importer establishes that a valid export certificate is in effect for certain sugar-containing products subject to a tariff-rate quota, that are products of a participating country, as defined in regulations of the United States Trade Representative (USTR). The export certificate is necessary to enable the importer to claim the in-quota rate of duty on the sugar-containing products.

EFFECTIVE DATE: July 14, 2000. FOR FURTHER INFORMATION CONTACT: Leon Hayward, Office of Field

Operations, (202–927–9704).

SUPPLEMENTARY INFORMATION:

Background

As a result of the Uruguay Round Agreements, approved by Congress in section 101 of the Uruguay Round Agreements Act (URAA) (Pub. L. 103– 465), the President, by Presidential Proclamation No. 6763, established a tariff-rate quota for imported sugarcontaining products.

Under a tariff-rate quota, the United States applies one tariff rate, known as the in-quota tariff rate, to imports of a product up to a particular amount, known as the in-quota quantity, and another, higher rate, known as the overquota rate, to imports of a product in excess of the given amount. The preferential, in-quota tariff rate would be applicable only to the extent that the aggregate in-quota quantity of a product allocated to a country had not been exceeded.

Under Presidential Proclamation No. 7235, dated October 7, 1999, the United States Trade Representative (USTR) was given authority under section 404(a) of the URAA to implement the tariff-rate quota for sugar-containing products to ensure that they do not disrupt the

orderly marketing of such products in the United States. The USTR has already assigned Canada an in-quota allocation of the sugar-containing products (64 FR 54719; October 7, 1999).

As part of the implementation of this tariff-rate quota, the USTR has established an export-certificate program under which exporting countries that have an allocation of the in-quota quantity and that wish to participate in the program may use export certificates for their sugarcontaining products that are exported to the United States. The USTR issued regulations for this export-certificate program (15 CFR part 2015) (64 FR 67152; December 1, 1999).

An exporting country wishing to participate in the export-certificate program must notify the USTR and provide the necessary supporting information. As defined in the USTR regulations (15 CFR 2015.2(e)), a participating country is a country that has received an allocation of the inquota quantity of the tariff-rate quota, and that the USTR has determined, and has so informed Customs, is eligible to use export certificates for their sugarcontaining products exported to the United States. The USTR has stated that it intends to publish a notice in the Federal Register whenever a country becomes, or ceases to be, a participating

The particular sugar-containing products subject to a tariff-rate quota for which the USTR has established the export-certificate program are described in additional U.S. Note 8 to chapter 17 of the Harmonized Tariff Schedule of the United States (HTSUS). Specifically, unless excepted as provided in additional U.S. Note 3 to chapter 17, HTSUS, the imported sugar-containing products covered by the exportcertificate program contain over 10 percent by dry weight of sugars derived from cane or sugar beets, whether or not mixed with other ingredients, and they are classified under one of the following HTSUS subheadings: 1701.91.54, 1704.90.74, 1806.20.75, 1806.20.95, 1806.90.55, 1901.90.56, 2101.12.54, 2101.20.54, 2106.90.78, or 2106.90.95.

While a country does not need to participate in the export-certificate program in order to receive the in-quota tariff rate for its share of the in-quota quantity, using export certificates assures the exporting country that only those exported sugar-containing products that it intends for the United States market are counted against its inquota allocation. As already noted, this helps ensure that such products do not disrupt the orderly marketing of sugar-

containing products in the United States.

On December 4, 1998, the Governments of the United States and Canada entered into a Record of Understanding regarding Areas of Agricultural Trade. In Annex 17 of this Record of Understanding, the United States agreed to require an export permit issued by the Government of Canada in order to enable an importer to claim the in-quota tariff rate for those sugarcontaining products of Canadian origin described in additional U.S. Note 8 to chapter 17, HTSUS. Canada was thus a participating country in this exportcertificate program as of January 31, 2000, the effective date of the USTR rule.

In accordance with the rulemaking of the USTR, by a document published in the Federal Register (65 FR 5430) on February 4, 2000, Customs issued an interim rule that added a new § 132.17 to the Customs Regulations (19 CFR 132.17), in order to prescribe the form and manner by which an importer establishes that a valid export certificate exists, including a unique number for the certificate that must be referenced on the entry or withdrawal from warehouse for consumption, whether filed in paper form or electronically. This was intended to ensure that no imports of the specified sugarcontaining products of a participating country are counted against the country's in-quota allocation unless the products are covered by a proper export certificate. The export certificate is necessary in this regard in order to enable the importer to claim the inquota rate of duty on the sugarcontaining products.

In addition, the Interim (a)(1)(A) List set forth as an Appendix to part 163, Customs Regulations (19 CFR part 163, Appendix), that lists the records required for the entry of merchandise, was revised to add a reference to the requirement in new § 132.17 that an importer possess a valid export certificate for sugar-containing products that are subject to a tariff-rate quota and that are products of a participating country, in order for the importer to be able to claim the applicable in-quota rate of duty.

Also, § 132.15, Customs Regulations (19 CFR 132.15), was revised to make provision for electronic entry filing in the case of beef subject to a tariff-rate quota, for which the importer must similarly possess a valid export certificate in order to claim the in-quota rate of duty.

No comments were received from the public in response to the interim rule, and Customs has now determined to 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,

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,

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

2000.

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