

**DEPARTMENT OF COMMERCE****Bureau of Industry and Security****15 CFR Part 745**

[Docket No. 061101286–7039–02]

RIN 0694–AD85

**Revisions to the Export Administration Regulations Based on U.S. Recognition of Montenegro as a Sovereign State; Correction****AGENCY:** Bureau of Industry and Security, Commerce.**ACTION:** Correcting amendment.

**SUMMARY:** The Bureau of Industry and Security (BIS) published a final rule in the *Federal Register* on Monday, November 27, 2006 (71 FR 68438) that amended the Export Administration Regulations (EAR) to add “Montenegro” and “Serbia” as separate countries in the EAR and to establish separate export licensing requirements for Montenegro and Serbia. The November 27, 2006, final rule omitted a conforming amendment to the list of States Parties to the Chemical Weapons Convention in the EAR. This document corrects that rule by listing Serbia and Montenegro as separate countries on that list.

**DATES:** *Effective Date:* This rule is effective March 19, 2007.

**ADDRESSES:** Although this is a final rule, comments are welcome and should be sent to [publiccomments@bis.doc.gov](mailto:publiccomments@bis.doc.gov), fax (202) 482–3355, or to Regulatory Policy Division, Bureau of Industry and Security, Room H2705, U.S. Department of Commerce, Washington, DC 20230. Please refer to regulatory identification number (RIN) 0694–AD85 in all comments, and in the subject line of e-mail comments. Comments on the collection of information should be sent to David Rostker, Office of Management and Budget (OMB), by e-mail to [David\\_Rostker@omb.eop.gov](mailto:David_Rostker@omb.eop.gov), or by fax to (202) 395–7285.

**FOR FURTHER INFORMATION CONTACT:** Steven Emme, Regulatory Policy Division, Bureau of Industry and Security, *Telephone:* (202) 482–2440.

**SUPPLEMENTARY INFORMATION:****Background**

This document corrects an inadvertent omission in the final rule that was published by the Bureau of Industry and Security (BIS) on November 27, 2006 (71 FR 68438). The November 27, 2006 final rule did not include a conforming amendment to Supplement No. 2 to part 745 of the EAR to amend the reference to “Serbia and Montenegro”. This document

corrects Supplement No. 2 to part 745 by removing “Serbia and Montenegro” and adding “Montenegro” and “Serbia”.

Although the Export Administration Act expired on August 20, 2001, the President, through Executive Order 13222 of August 17, 2001, 3 CFR, 2001 Comp., p. 783 (2002), as extended by the Notice of August 3, 2006, 71 FR 44551 (August 7, 2006), has continued the Export Administration Regulations in effect under the International Emergency Economic Powers Act.

**Rulemaking Requirements**

1. This final rule has been determined to be not significant for purposes of E.O. 12866.

2. Notwithstanding any other provision of law, no person is required to respond to, nor shall any person be subject to a penalty for failure to comply with, a collection of information subject to the requirements of the Paperwork Reduction Act of 1995 (44 U.S.C. 3501 *et seq.*) (PRA), unless that collection of information displays a currently valid Office of Management and Budget (OMB) Control Number. This rule contains a collection of information subject to the requirements of the PRA. This collection has previously been approved by OMB under control number 0694–0088 (Multi-Purpose Application), which carries a burden hour estimate of 58 minutes to prepare and submit form BIS–748. This rule is not expected to result in any change for collection purposes.

3. This rule does not contain policies with Federalism implications as that term is defined under E.O. 13132.

4. The provisions of the Administrative Procedure Act (5 U.S.C. 553) requiring notice of proposed rulemaking, the opportunity for public participation, and a delay in effective date, are inapplicable because this regulation involves a military and foreign affairs function of the United States (5 U.S.C. 553(a)(1)). Further, no other law requires that a notice of proposed rulemaking and an opportunity for public comment be given for this final rule. Because a notice of proposed rulemaking and an opportunity for public comment are not required to be given for this rule under the Administrative Procedure Act or by any other law, the analytical requirements of the Regulatory Flexibility Act (5 U.S.C. 601 *et seq.*) are not applicable. 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 Steven Emme, Regulatory Policy

Division, Bureau of Industry and Security, Department of Commerce, P.O. Box 273, Washington, DC 20044.

**List of Subjects in 15 CFR Part 745**

Administrative practice and procedure, Chemicals, Exports, Foreign trade, Reporting and recordkeeping requirements.

■ Accordingly, part 745 of the Export Administration Regulations (15 CFR parts 730–799) is corrected by making the following correcting amendment:

**PART 745—[AMENDED]**

■ 1. The authority citation for 15 CFR part 745 continues to read as follows:

**Authority:** 50 U.S.C. 1701 *et seq.*; E.O. 12938, 59 FR 59099, 3 CFR, 1994 Comp., p. 950; Notice of October 27, 2006, 71 FR 64109 (October 31, 2006).

■ 2. In Supplement No. 2 to part 745, States Parties to the Convention on the Prohibition of the Development, Production, Stockpiling, and Use of Chemical Weapons and on Their Destruction is amended by removing “Serbia and Montenegro” and by adding, in alphabetical order, “Montenegro” and “Serbia”.

Dated: March 9, 2007.

**Eileen Albanese,**

*Director, Office of Exporter Services.*

[FR Doc. 07–1275 Filed 3–16–07; 8:45 am]

**BILLING CODE 3510–33–P**

**DEPARTMENT OF COMMERCE****National Oceanic and Atmospheric Administration****15 CFR Part 922**

[Docket No. 031001243–6227–02]

RIN 0648–AQ41

**Gray’s Reef National Marine Sanctuary Regulations; Announcement of Effective Date**

**AGENCY:** National Marine Sanctuary Program (NMSP), National Ocean Service (NOS), National Oceanic and Atmospheric Administration (NOAA), Department of Commerce (DOC).

**ACTION:** Announcement of Effective Date.

**SUMMARY:** On October 12, 2006, the National Oceanic and Atmospheric Administration (NOAA) published a final rule (71 FR 60055) issuing a final management plan, revised designation document and final regulations for the Gray’s Reef National Marine Sanctuary. Under the National Marine Sanctuaries Act, the final regulations would

automatically take effect at the end of 45 days of continuous session of Congress beginning on October 12, 2006. The 45-day review period ended on February 16, 2007. This document confirms the effective date as February 16, 2007.

**DATES:** *Effective Date:* The final rule published on October 12, 2006 (71 FR 60055) took effect on February 16, 2007.

**FOR FURTHER INFORMATION CONTACT:** Becky Shortland, Gray's Reef National Marine Sanctuary, 10 Ocean Science Circle, Savannah, Georgia 31411; 912-598-2381; [Becky.Shortland@noaa.gov](mailto:Becky.Shortland@noaa.gov). (Federal Domestic Assistance Catalog Number 11.429 Marine Sanctuary Program) Dated: March 13, 2007.

**John H. Dunnigan,**

*Assistant Administrator for Ocean Services and Coastal Zone Management.*

[FR Doc. 07-1303 Filed 3-16-07; 8:45 am]

**BILLING CODE 3510-08-M**

## SOCIAL SECURITY ADMINISTRATION

### 20 CFR Part 416

#### Revised Medical Criteria for Determination of Disability, Musculoskeletal System and Related Criteria

##### *CFR Correction*

In Title 20 of the Code of Federal Regulations, Parts 400 to 499, revised as of April 1, 2006, on page 948, § 416.933 is corrected by adding a sentence after the second sentence to read as follows:

#### **§ 416.933 How we make a finding of presumptive disability or presumptive blindness.**

\* \* \* For other impairments, a finding of disability or blindness must be based on medical evidence or other information that, though not sufficient for a formal determination of disability or blindness, is sufficient for us to find that there is a high degree of probability that you are disabled or blind. \* \* \*

[FR Doc. 07-55503 Filed 3-16-07; 8:45 am]

**BILLING CODE 1505-01-D**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

#### 21 CFR Part 341

[Docket No. 1976N-0052G] (formerly Docket No. 76N-052G)

#### Cold, Cough, Allergy, Bronchodilator, and Antiasthmatic Drug Products for Over-the-Counter Human Use; Technical Amendment

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Final rule; technical amendment.

**SUMMARY:** The Food and Drug Administration (FDA) is amending its regulations to change the location of a section in an over-the-counter (OTC) drug monograph. This action is editorial in nature and is intended to improve the accuracy of the agency's regulations.

**DATES:** This rule is effective March 19, 2007.

**FOR FURTHER INFORMATION CONTACT:** Gerald M. Rachanow, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 22, rm. 5496, Silver Spring, MD 20993, 301-796-2090.

**SUPPLEMENTARY INFORMATION:** FDA published the final monograph (FM) for cold, cough, allergy, bronchodilator, and antiasthmatic combination drug products for OTC human use in the **Federal Register** of December 23, 2002 (67 FR 78158). In that FM, FDA inadvertently added § 341.40 (21 CFR 341.40) to subpart C of the monograph, when that section should have been added to subpart B of the monograph. Accordingly, FDA is now moving § 341.40 from subpart C to subpart B of the monograph.

Publication of this document constitutes final action on this change under the Administrative Procedure Act (5 U.S.C. 553). Notice and public procedures are unnecessary because FDA is merely implementing a change in the location of a section in an OTC drug monograph. No other changes are being made to that section of the monograph.

#### List of Subjects in 21 CFR Part 341

Labeling, Over-the-counter drugs.

■ Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR part 341 is amended as follows:

## PART 341—COLD, COUGH, ALLERGY, BRONCHODILATOR, AND ANTI-ASTHMATIC DRUG PRODUCTS FOR OVER-THE-COUNTER HUMAN USE

■ 1. The authority citation for 21 CFR part 341 continues to read as follows:

**Authority:** 21 U.S.C. 321, 351, 352, 353, 355, 360, 371.

#### Subpart B—Active Ingredients [Amended]

■ 2. Remove § 341.40 *Permitted combinations of active ingredients* from subpart C and add it to subpart B of part 341.

Dated: March 12, 2007.

**Jeffrey Shuren,**

*Assistant Commissioner for Policy.*

[FR Doc. E7-4957 Filed 3-16-07; 8:45 am]

**BILLING CODE 4160-01-S**

## DEPARTMENT OF THE TREASURY

### 31 CFR Part 103

RIN 1506-AA83

#### Financial Crimes Enforcement Network; Amendment to the Bank Secrecy Act Regulations—Imposition of Special Measure Against Banco Delta Asia, Including Its Subsidiaries Delta Asia Credit Limited and Delta Asia Insurance Limited, as a Financial Institution of Primary Money Laundering Concern

**AGENCY:** Financial Crimes Enforcement Network, Department of the Treasury.

**ACTION:** Final rule.

**SUMMARY:** The Financial Crimes Enforcement Network ("FinCEN") is issuing a final rule imposing a special measure against Banco Delta Asia SARL ("Banco Delta Asia" or "the bank") as a financial institution of primary money laundering concern, pursuant to the authority contained in 31 U.S.C. 5318A of the Bank Secrecy Act.

**DATES:** This final rule is effective on April 18, 2007.

**FOR FURTHER INFORMATION CONTACT:** Regulatory Policy and Programs Division, Financial Crimes Enforcement Network, (800) 949-2732.

#### SUPPLEMENTARY INFORMATION:

##### I. Background

##### A. Statutory Provisions

On October 26, 2001, the President signed into law the Uniting and Strengthening America by Providing Appropriate Tools Required To