SUPPLEMENT NO. 4 TO PART 744—ENTITY LIST—Continued

Country	Entity	License requirement	License review policy	**Federal Register** citation
	#701 Damas Tower, 702 Al Maktoum St., Dubai, U.A.E.; *and* 701 Attar Tower, Maktoum St. Dubai, U.A.E.; *and* City Tower, Al Maktoum St., Office No. 701, Dubai U.A.E.; *and* P.O. Box 40595, Dubai, U.A.E.; *and* Warehouse No. 8, Plot No. 238, Rashidiya, Dubai, U.A.E..			
	* *	*	*	*
	Rainbow General Trading Company, ... City Tower 2, 20th Floor, Office #2005, Sheikh Zayed Road, Dubai, U.A.E..	For all items subject to the EAR. (See § 744.11 of the EAR)	Presumption of denial	81 FR [INSERT FR PAGE NUMBER], 2/23/16.
	* *	*	*	*
UNITED KINGDOM	* *	*	*	*
	Veteran Avia LLC a.k.a., the following alias:. —Veteran Airline 1 Beckett Place, South Hamptonshire, London, U.K. (See also addresses under Armenia, Greece, India, and Pakistan)..	For all items subject to the EAR. (See § 744.11 of the EAR)	Presumption of denial	79 FR 56003, 9/18/14. 81 FR [INSERT FR PAGE NUMBER], 2/23/16.
	* *	*	*	*

Dated: February 16, 2016.

Kevin J. Wolf,

Assistant Secretary for Export Administration.

[FR Doc. 2016–03745 Filed 2–22–16; 8:45 am]

BILLING CODE 3510–33–P

JOINT BOARD FOR THE ENROLLMENT OF ACTUARIES

20 CFR Part 900

[TD 9749]

RIN 1545–BM81

Regulations Governing Organization of the Joint Board for the Enrollment of Actuaries

AGENCY: Joint Board for the Enrollment of Actuaries.

ACTION: Final rule.

SUMMARY: This document contains final regulations relating to the organization of the Joint Board for the Enrollment of Actuaries. The regulations are being amended in order to conform one provision of the regulations to the Bylaws of the Joint Board. These regulations solely address the internal management of the Joint Board and do not affect pension plans, plan participants, actuaries, or the general public.

DATES: *Effective date:* These regulations are effective *April 25, 2016.*

FOR FURTHER INFORMATION CONTACT: Patrick McDonough, Executive Director, Joint Board for the Enrollment of Actuaries, at (703) 414–2173 (not a toll-free number).

SUPPLEMENTARY INFORMATION:

Background and Explanation

The Joint Board for the Enrollment of Actuaries was established on October 31, 1974 pursuant to section 3041 of the Employee Retirement Income Security Act of 1974 (88 Stat. 829), Public Law 93–406 (ERISA). Section 3041 of ERISA provides that the Secretary of Labor and the Secretary of the Treasury shall, not later than the last day of the first calendar month beginning after the date of enactment of ERISA, establish a Joint Board for the Enrollment of Actuaries (Joint Board).

Regulations under ERISA section 3041 were published in the **Federal Register** on April 30, 1975 (40 FR 18776) and are currently located in the Code of Federal Regulations at 20 CFR part 900 (the 1975 Joint Board regulations). These regulations provide that, pursuant to the Bylaws, three members are appointed by the Secretary of the Treasury, two members are appointed by the Secretary of Labor, the Chairman of the Joint Board is to be elected from among the Treasury Department representatives, and the Secretary is to be elected from among the Labor Department representatives.

On April 27, 1981, the Secretaries of Treasury and Labor approved restated Bylaws of the Joint Board (the 1981 Bylaws). Sections 3(b) and 3(c) of the 1981 Bylaws provide that the Chairman and Secretary, respectively, will be elected for a one-year term by the Joint Board from among its members, eliminating the requirement that the Chairman be a Treasury Department representative and the Secretary be a Labor Department representative.

These final regulations amend § 900.3 of the 1975 Joint Board regulations in order to conform the regulations to the 1981 Bylaws.

Special Analyses

These regulations are being published as a final rule because the amendments apply solely to the Joint Board's organization and management. Moreover, the Joint Board finds good cause that these changes do not impose any requirements on any member of the public. These amendments are the most efficient means for the Joint Board to harmonize the regulations and Bylaws involving the Board's internal election procedure.

Accordingly, pursuant to 5 U.S.C. 553(a)(2), 553(b)(3)(A), and 553(b)(3)(B), the Joint Board finds good cause that prior notice and other public procedures with respect to this rule are unnecessary. Because a notice of proposed rulemaking is not required, the provisions of the Regulatory Flexibility Act, 5 U.S.C. 601–612, do not apply.

This rule is not a significant regulatory action pursuant to Executive Order 12866, as supplemented and reaffirmed by Executive Order 13563. Therefore, a regulatory impact assessment is not required.

List of Subjects in 20 CFR Part 900

Organization and functions (Government agencies).

Adoption of Amendments to the Regulations

Accordingly, 20 CFR part 900 is amended as follows:

PART 900—STATEMENT OF ORGANIZATION

■ **Paragraph 1.** The authority citation for part 900 continues to read as follows:

Authority: Sec. 3041–2, Pub. L. 93–406, 88 Stat. 829, 1002 (29 U.S.C. 1241–2).

■ **Par. 2.** Section 900.3 is revised to read as follows:

§ 900.3 Composition.

Pursuant to the Bylaws, the Joint Board consists of three members appointed by the Secretary of the Treasury and two members appointed by the Secretary of Labor. The Board elects a Chairman and a Secretary from among the Department of the Treasury and the Department of Labor members. The Pension Benefit Guaranty Corporation may designate a non-voting representative to sit with, and participate in, the discussions of the Board. All decisions of the Board are made by simple majority vote.

Approved: February 12, 2016.

Carolyn E. Zimmerman,

Chairman, Joint Board for the Enrollment of Actuaries.

[FR Doc. 2016–03655 Filed 2–22–16; 8:45 am]

BILLING CODE 4830–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 101

[Docket No. FDA–2016–N–0585]

Food Labeling: Nutrient Content Claims; Alpha-Linolenic Acid, Eicosapentaenoic Acid, and Docosahexaenoic Acid Omega-3 Fatty Acids; Small Entity Compliance Guide; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification of availability.

SUMMARY: The Food and Drug Administration (FDA, the Agency, or we) is announcing the availability of a guidance for industry entitled ''Food Labeling: Nutrient Content Claims; Alpha-Linolenic Acid, Eicosapentaenoic Acid, and Docosahexaenoic Acid Omega-3 Fatty Acids—Small Entity Compliance Guide.'' The small entity compliance guide (SECG) is intended to help small entities comply with the final rule titled ''Food Labeling: Nutrient Content Claims; Alpha-Linolenic Acid, Eicosapentaenoic Acid, and Docosahexaenoic Acid Omega-3 Fatty Acids.''

DATES: Submit either electronic or written comments on FDA guidances at any time.

ADDRESSES: You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: http://www.regulations.gov*. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *http://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *http://www.regulations.gov*.
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand delivery/Courier (for written/paper submissions):* Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Division of Dockets Management, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2016–N–0585 for ''Food Labeling: Nutrient Content Claims; Alpha-Linolenic Acid, Eicosapentaenoic Acid, and Docosahexaenoic Acid Omega-3 Fatty Acids; Small Entity Compliance Guide.'' Received comments will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *http://www.regulations.gov* or at the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

- Confidential Submissions—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *http://www.regulations.gov*. Submit both copies to the Division of Dockets Management. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *http://www.fda.gov/regulatoryinformation/dockets/default.htm*.

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *http://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Division of Dockets Management, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

Submit written requests for single copies of the SECG to the Office of Nutrition and Food Labeling, Center for Food Safety and Applied Nutrition (HFS–830), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740. Send two self-addressed adhesive labels to assist that office in processing your request. See the **SUPPLEMENTARY**