form, with a request for confidential treatment, and you have to follow the procedure explained in FTC Rule 4.9(c), 16 CFR 4.9(c). Your comment will be kept confidential only if the FTC General Counsel, in his or her sole discretion, grants your request in accordance with the law and the public interest.

Postal mail addressed to the Commission is subject to delay due to heightened security screening. As a result, we encourage you to submit your comments online, or to send them to the Commission by courier or overnight service. To make sure that the Commission considers your online comment, you must file it at *https://ftcpublic.commentworks.com/ftc/altfuelspra2,* by following the instructions on the web-based form. If this Notice appears at *http://www.regulations.gov/#!home,* you also may file a comment through that Web site.

If you file your comment on paper, write ''Paperwork Comment: FTC File No. P134200'' on your comment and on the envelope, and mail or deliver it to the following address: Federal Trade Commission, Office of the Secretary, Room H–113 (Annex J), 600 Pennsylvania Avenue NW., Washington, DC 20580. If possible, submit your paper comment to the Commission by courier or overnight service.

Visit the Commission Web site at *http://www.ftc.gov* to read this Notice. The FTC Act and other laws that the Commission administers permit the collection of public comments to consider and use in this proceeding as appropriate. The Commission will consider all timely and responsive public comments that it receives on or before April 4, 2013. You can find more information, including routine uses permitted by the Privacy Act, in the Commission's privacy policy, at *http://www.ftc.gov/ftc/privacy.htm.*

Comments on the information collection requirements subject to review under the PRA should additionally be submitted to OMB. If sent by U.S. mail, they should be addressed to Office of Information and Regulatory Affairs, Office of Management and Budget, Attention: Desk Officer for the Federal Trade Commission, New Executive Office Building, Docket Library, Room 10102, 725 17th Street NW., Washington, DC 20503. Comments sent to OMB by U.S. postal mail, however, are subject to delays due to heightened security precautions. Thus, comments instead should be sent by facsimile to (202) 395–5167.

David C. Shonka,

Acting General Counsel.

[FR Doc. 2013–05070 Filed 3–4–13; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Statement of Delegation of Authority; Health Resources and Services Administration and Centers for Disease Control and Prevention

I hereby delegate to the Administrator, Health Resources and Services Administration (HRSA), and the Director, Centers for Disease Control and Prevention (CDC), with authority to redelegate, the authority vested in the Secretary of the Department of Health and Human Services under Title III, Part R, Section 399BB, titled ''Autism, Education, Early Detection, and Intervention,'' of the Public Health Service (PHS) Act, as amended, insofar as such authority pertains to the functions of HRSA and CDC, respectively. HRSA and CDC will coordinate and collaborate with each other, with the National Institutes of Health (NIH), and with the Administration for Children and Families (ACF), as appropriate, in implementing this authority. In addition, nothing in this delegation of authority would preclude NIH from pursuing research and training activities authorized by the PHS Act. HRSA and CDC will also coordinate and collaborate with other agencies, as appropriate, in implementing this authority.

This delegation excludes the authority to issue regulations, to establish advisory committees and councils and appoint their members, and shall be exercised in accordance with the Department's applicable policies, procedures, and guidelines.

I hereby affirm and ratify any actions taken by the Administrator, HRSA, the Director, CDC, or other HRSA and CDC officials, which involve the exercise of these authorities prior to the effective date of this delegation.

This delegation is effective upon date of signature.

Dated: February 22, 2013.

Kathleen Sebelius,

Secretary.

[FR Doc. 2013–04946 Filed 3–4–13; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Notice of Interest Rate on Overdue Debts

Section 30.18 of the Department of Health and Human Services' claims collection regulations (45 CFR part 30) provides that the Secretary shall charge an annual rate of interest, which is determined and fixed by the Secretary of the Treasury after considering private consumer rates of interest on the date that the Department of Health and Human Services becomes entitled to recovery. The rate cannot be lower than the Department of Treasury's current value of funds rate or the applicable rate determined from the ''Schedule of Certified Interest Rates with Range of Maturities'' unless the Secretary waives interest in whole or part, or a different rate is prescribed by statute, contract, or repayment agreement. The Secretary of the Treasury may revise this rate quarterly. The Department of Health and Human Services publishes this rate in the **Federal Register**.

The current rate of 10⅝%, as fixed by the Secretary of the Treasury, is certified for the quarter ended December 31, 2012. This interest rate is effective until the Secretary of the Treasury notifies the Department of Health and Human Services of any change.

Dated: February 25, 2013.

Margie Yanchuk,

Director, Office of Financial Policy and Reporting.

[FR Doc. 2013–04945 Filed 3–4–13; 8:45 am]

BILLING CODE 4150–04–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Subcommittee for Dose Reconstruction Reviews (SDRR), Advisory Board on Radiation and Worker Health (ABRWH or the Advisory Board), National Institute for Occupational Safety and Health (NIOSH)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC), announces the following meeting for the aforementioned subcommittee:

Time and Date: 9:00 a.m.–5:00 p.m. Eastern Time, March 25, 2013.

Place: Cincinnati Airport Marriott, 2395 Progress Drive, Hebron, Kentucky

41018, Telephone (859) 334–4611, Fax (859) 334–4619.

Status: Open to the public, but without an oral public comment period.

The USA toll-free, dial-in number is 1–866–659–0537 with a pass code of 9933701.

Background: The Advisory Board was established under the Energy Employees Occupational Illness Compensation Program Act of 2000 to advise the President on a variety of policy and technical functions required to implement and effectively manage the new compensation program. Key functions of the Advisory Board include providing advice on the development of probability of causation guidelines that have been promulgated by the Department of Health and Human Services (HHS) as a final rule; advice on methods of dose reconstruction, which have also been promulgated by HHS as a final rule; advice on the scientific validity and quality of dose estimation and reconstruction efforts being performed for purposes of the compensation program; and advice on petitions to add classes of workers to the Special Exposure Cohort (SEC).

In December 2000, the President delegated responsibility for funding, staffing, and operating the Advisory Board to HHS, which subsequently delegated this authority to CDC. NIOSH implements this responsibility for CDC. The charter was issued on August 3, 2001, renewed at appropriate intervals, and will expire on August 3, 2013.

Purpose: The Advisory Board is charged with (a) providing advice to the Secretary, HHS, on the development of guidelines under Executive Order 13179; (b) providing advice to the Secretary, HHS, on the scientific validity and quality of dose reconstruction efforts performed for this program; and (c) upon request by the Secretary, HHS, advise the Secretary on whether there is a class of employees at any Department of Energy facility who were exposed to radiation but for whom it is not feasible to estimate their radiation dose, and on whether there is reasonable likelihood that such radiation doses may have endangered the health of members of this class. The Subcommittee for Dose Reconstruction Reviews was established to aid the Advisory Board in carrying out its duty to advise the Secretary, HHS, on dose reconstruction.

Matters To Be Discussed: The agenda for the Subcommittee meeting includes: dose reconstruction program quality management and assurance activities, including: Current findings from NIOSH internal dose reconstruction blind reviews; discussion of dose reconstruction cases under review (sets 8–9, Savannah River Site, Rocky Flats Plant, and Los Alamos National Laboratory cases from sets 10–13); and selection of ''blind dose reconstructions'' from set 16.

The agenda is subject to change as priorities dictate.

In the event an individual cannot attend, written comments may be submitted. Any written comments received will be provided at the meeting and should be submitted to the contact person below well in advance of the meeting.

Contact Person for More Information: Theodore Katz, Designated Federal Official, NIOSH, CDC, 1600 Clifton Road, Mailstop E–20, Atlanta, Georgia 30333, Telephone (513) 533–6800, Toll Free 1 (800) CDC–INFO, email *ocas@cdc.gov*.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dana Redford,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2013–04969 Filed 3–4–13; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0777]

Adrian Vela: Debarment Order

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is issuing an order under the Federal Food, Drug, and Cosmetic Act (the FD&C Act) debarring Adrian Vela for a period of 5 years from importing articles of food or offering such articles for importation into the United States. FDA bases this order on a finding that Mr. Vela was convicted of three felony counts under Federal law for conduct relating to the importation into the United States of an article of food. Mr. Vela was given notice of the proposed debarment and an opportunity to request a hearing within the timeframe prescribed by regulation. As of November 3, 2012 (30 days after receipt of the notice), Mr. Vela had not responded. Mr. Vela's failure to respond constitutes a waiver of his right to a hearing concerning this action.

DATES: This order is effective March 5, 2013.

ADDRESSES: Submit applications for termination of debarment to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Kenny Shade, Office of Regulatory Affairs, Food and Drug Administration, 12420 Parklawn Dr., Rockville, MD 20857, 301–796–4640.

SUPPLEMENTARY INFORMATION:

I. Background

Section 306(b)(1)(C) of the FD&C Act (21 U.S.C. 335a(b)(1)(C)) permits FDA to debar an individual from importing an article of food or offering such an article for import into the United States if FDA finds, as required by section 306(b)(3)(A) of the FD&C Act, that the individual has been convicted of a felony for conduct relating to the importation into the United States of any food.

On November 21, 2011, Mr. Vela was convicted, as defined in section 306(l)(1)(B) of the FD&C Act, when the U.S. District Court for the Southern District of Florida accepted his plea of guilty and entered judgment against him for the following offenses: One count of conspiracy to falsely label and misbrand food, in violation of 18 U.S.C. 371; one count of false labeling of seafood under the Lacey Act, in violation of 16 U.S.C. 3372(d)(2); and one count of misbranding food in violation of 21 U.S.C. 331(a).

FDA's finding that debarment is appropriate is based on the felony convictions referenced herein for conduct relating to the importation into the United States of any food. The factual basis for these convictions is as follows: As alleged in the criminal information filed against Mr. Vela, he was the operating manager and sole shareholder of Sea Food Center, a seafood wholesaler engaged in various aspects of purchasing, importing, processing, packing, selling, and exporting seafood products.

Beginning on or about June 30, 2008, and continuing through on or about June 29, 2009, he knowingly combined, conspired, confederated, and agreed with his co-conspirators to commit an offense against the laws of the United States related to the importation of food. The purpose of the conspiracy was for Mr. Vela and his co-conspirators to unlawfully enrich themselves by