Federal Officer, NIOSH, CDC, 1600 Clifton Road, Mailstop E–20, Atlanta, Georgia 30329, Telephone (513) 533–6800, Toll Free 1 (800) CDC–INFO, Email *ocas@cdc.gov.*

SUPPLEMENTARY INFORMATION:

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, rechartered on February 12, 2018, pursuant to Executive Order 13708, and will terminate on September 30, 2019.

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. SPR is responsible for overseeing, tracking, and participating in the reviews of all procedures used in the dose reconstruction process by the NIOSH Division of Compensation Analysis and Support (DCAS) and its dose reconstruction contractor (Oak Ridge Associated Universities—ORAU).

Matters to be Considered: The agenda will include discussions on the following dose reconstruction procedures: (a) Procedures associated specifically with the following sites: Norton Company, Paducah, Blockson Chemical Company, DuPont Deepwater Works, Huntington Pilot Plan, Y–12, Aliquippa Forge, Hooker Electrochemical Plant; (b) procedures associated with Atomic Weapons Employers generally; and, (c) general procedures for dose reconstructions. Agenda items are subject to change as priorities dictate.

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.

Sherri Berger,

Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2018–20016 Filed 9–13–18; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—CE19–001, Injury Control Research Centers; Amended Notice of Meeting

Notice is hereby given of a change in the meeting of the Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—CE19–001, Injury Control Research Centers; October 30 and November 2, 2018, 8:30 a.m.–5:00 p.m., EDT, in the original FRN.

The Georgian Terrace, 659 Peachtree St. NE, Atlanta, GA 30308 which was published in the **Federal Register** on August 23, 2018, Volume 83, Number 164, pages 42655–42656.

The meeting is being amended to change the location and time to the Sheraton Atlanta Hotel, 165 Courtland Street NE, Atlanta, GA 30303; October 30–November 2, 2018, 8:00 a.m.–5:30 p.m., EDT. The meeting is closed to the public.

FOR FURTHER INFORMATION CONTACT: Mikel L. Walters, M.A., Ph.D., Scientific Review Official, NCIPC, CDC, 4770 Buford Highway NE, Mailstop F–63, Atlanta, Georgia 30341, (404) 639–0913; *mwalters@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.

Sherri Berger,

Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2018–20024 Filed 9–13–18; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Intent To Award a Single-Source Supplement; Notice

ACTION: Announcing the Intent to Award a Single-Source Supplement to provide the National Aging Network with timely, relevant, high quality opportunities to further enhance their knowledge and skills related to nutrition services.

SUMMARY: The Administration for Community Living (ACL) announces the intent to award a single-source supplement to the current cooperative agreement held by Meals on Wheels America for the project *Enhancing the Knowledge and Skills of the Aging Network.*

FOR FURTHER INFORMATION CONTACT: For further information or comments regarding this program supplement, contact Keri Lipperini, U.S. Department of Health and Human Services, Administration for Community Living, Administration on Aging, Office of Nutrition and Health Promotion Programs, 202–795–7422, email *keri.lipperini@acl.hhs.gov.*

SUPPLEMENTARY INFORMATION: The purpose of this supplement is to: (1) Support the development and dissemination of resources for experienced and inexperienced Aging Network Nutrition Program providers; and (2) enhance peer-learning opportunities for State Units on Aging (SUAs), Area Agencies on Aging (AAAs), and Nutrition Program providers.

The administrative supplement for FY 2018 will be in the amount of $175,242, bringing the total award for FY 2018 to $400,001.

The additional funding will not be used to begin new projects, but it will be used to enhance existing efforts. The grantee will continue to provide appropriate, quality nutrition-related

resources, address new opportunities to embed nutrition services within the home and community-based service systems, and engage successfully in emerging models of integrated health care.

Program Name: Enhancing the Knowledge and Skills of the Aging Network.

Recipient: Meals on Wheels America.

Period of Performance: The supplement award will be issued for the second year of a three year project period of Sept 1, 2017 to August 31, 2020.

Total Award Amount: $400,001 in FY 2018.

Award Type: Cooperative Agreement Supplement.

Statutory Authority: The Older Americans Act (OAA) of 1965, as amended, Public Law 114–144.

Basis for Award: Meals on Wheels America (MOWA) is currently funded to carry out the objectives of this project through its current project entitled, *National Resource Center on Nutrition and Aging* for the period of September 1, 2017 through August 31, 2020. Since the project's implementation, the grantee has made satisfactory progress toward its approved work plan. The supplement will enable the grantee to carry their work even further, enhancing the support they provide to the Aging Network Nutrition Program Providers. The additional funding will not be used to begin new projects or activities, but rather to enhance efforts specific to tribal populations and congregate meal settings.

MOWA is uniquely positioned to complete the work called for under this project. They have an already established infrastructure and are a known and trusted organization in the Aging Network. Prior to this current award, MOWA competed and was awarded the *National Nutrition Center* for 6 years. They have an established presence within much of the Aging Network. Under this current award period, they are providing educational opportunities for the Aging Network Nutrition Program Providers, including webinars and live trainings. They have a comprehensive, interactive web-based repository (*www.nutritionandaging.org*) with tools and resources, including—but not limited to—issues briefs, policy and practice models, and toolkits. They have also presented to the Aging Network locally and on a national level. They have reached thousands of providers using their: (1) Comprehensive database of SUAs, AAAs, and other Nutrition Program Providers; and (2) Leadership Academy, which provides expert consultation around nutrition program delivery and the use of technology to enhance services. In addition, they have developed partnerships with organizations, universities, and other entities to provide education and support for the Aging Network.

Establishing an entirely new grant project at this time would be potentially disruptive to the current work already well under way. More importantly, it could cause confusion among the Aging Network Nutrition Program Providers, which could have a negative effect on training and support opportunities. If this supplement were not provided, the project would be unable to address the significant unmet educational needs of the Aging Network Nutrition Program Providers.

Dated: September 5, 2018.

Mary Lazare,

Principal Deputy Administrator.

[FR Doc. 2018–19925 Filed 9–13–18; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–D–0456]

Appropriate Use of Voluntary Consensus Standards in Premarket Submissions for Medical Devices; Guidance for Industry and Food and Drug Administration Staff; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a final guidance entitled ''Appropriate Use of Voluntary Consensus Standards in Premarket Submissions for Medical Devices.'' Voluntary consensus standards can be a valuable resource for industry and FDA staff because such standards can increase predictability, streamline premarket review, provide clearer regulatory expectations, and facilitate market entry for safe and effective medical products. FDA developed this document to provide guidance to industry and FDA reviewers about the appropriate use of voluntary consensus standards in the preparation and evaluation of premarket submissions for medical devices. This guidance applies to all articles that meet the definition of a ''device'' under the Federal Food, Drug, and Cosmetic Act (FD&C Act).

DATES: The announcement of the guidance is published in the **Federal Register** on September 14, 2018.

ADDRESSES: You may submit either electronic or written comments on Agency guidances at any time as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://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 *https://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):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, 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–2014–D–0456 for ''Appropriate Use of Voluntary Consensus Standards in Premarket Submissions for Medical Devices.'' Received comments will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday.