

obligations, these statutory provisions provide the legal authorization for the collection of information on the FR 3016. The FR 3016 is a voluntary survey. No issue of confidentiality normally arises under the FR 3016, as names and any other characteristics that would permit personal identification of respondents are not reported to the Board. However, should the Board obtain such information, it would likely be exempt under exemption 6 of the Freedom of Information Act (5 U.S.C. 552(b)(6)) to the extent that it includes “personnel and medical files and similar files the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.”

Board of Governors of the Federal Reserve System, May 17, 2017.

Ann E. Misback,

Secretary of the Board.

[FR Doc. 2017-10331 Filed 5-19-17; 8:45 am]

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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: 10:30 a.m.–5:00 p.m., EDT, June 27, 2017.

Place: Audio Conference Call via FTS Conferencing.

Status: Open to the public, but without a public comment period. The public is welcome to submit written comments in advance of the meeting, to the contact person below. Written comments received in advance of the meeting will be included in the official record of the meeting. The public is also welcome to listen to the meeting by joining the teleconference at the USA toll-free, dial-in number at 1-866-659-0537 and the pass code is 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, rechartered on March 22, 2016 pursuant to Executive Order 13708, and will expire on September 30, 2017.

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 for Discussion: The agenda for the Subcommittee meeting includes the following dose reconstruction program quality management and assurance activities: Dose reconstruction cases under review from Sets 14–23, including the Oak Ridge sites (Y–12, K–25, Oak Ridge National Laboratory), Hanford, Feed Materials Production Center (“Fernald”), Mound Plant, Rocky Flats Plant, Nevada Test Site, Idaho National Laboratory, Savannah River Site, and other Department of Energy and “Atomic Weapons Employer” facilities.

The agenda is subject to change as priorities dictate.

Contact Person for More Information: Theodore Katz, Designated Federal

Officer, 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.

Elaine L. Baker,

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

[FR Doc. 2017-10332 Filed 5-19-17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention (CDC)

Board of Scientific Counselors, National Center for Environmental Health/Agency for Toxic Substances and Disease Registry (BSC, NCEH/ATSDR), Lead Poisoning Prevention (LPP) Subcommittee

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the CDC, National Center for Environmental Health/Agency for Toxic Substances and Disease Registry (NCEH/ATSDR) announces the following meeting of the aforementioned committee:

Time and date: 9:00 a.m.–12:00 p.m. EDT, June 23, 2017.

Place: The meeting will be accessible by teleconference. Please dial toll free 1-888-790-2009 Passcode: 7865774.

Status: Open to the public, via teleconference line. No limit on number of lines. The public is welcome to participate during the Public Comment period which is scheduled from 11:00 a.m. until 11:15 a.m. EST (15 minutes). Individuals wishing to make a comment during Public Comment period, please email your name, organization, and phone number by Monday, June 15, 2017 to Amanda Malasky at AMalasky@cdc.gov.

Purpose: The subcommittee will discuss strategies and options on ways to prioritize NCEH/ATSDR’s activities, improve health outcomes, and address health disparities as it relates to lead exposures. The subcommittee will deliberate on ways to evaluate lead exposure and how to best conduct health evaluations through exposure

and epidemiologic studies. Subcommittee proposals on lead poisoning prevention and national lead poisoning prevention efforts will be provided to the Board of Scientific Counselors for deliberation and possible adoption as formal recommendations to NCEH/ATSDR.

Matters for Discussion: Agenda items will include the following: Lead Poisoning Prevention Program (status), Flint Registry (status), Revision of Blood Lead Level reference value (status), Discussion of legislative requirements of a new Lead Exposure and Prevention Federal Advisory Committee, Federal partnership efforts.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Amanda Malasky, Coordinator, Lead Poisoning Prevention Subcommittee, BSC, NCEH/ATSDR, 4770 Buford Highway, Mail Stop F-45, Chamblee, Georgia 30345; telephone 770/488-7699, Fax: 770/488-3377; Email: AMalasky@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.

Elaine L. Baker,

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

[FR Doc. 2017-10333 Filed 5-19-17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Initial Review

The meeting announced below concerns the Centers for Disease Control and Prevention (CDC) initial review of applications in response to Funding Opportunity Announcement (FOA) GH16-006, Conducting Public Health Research in Kenya; GH17-004, Conducting Public Health Research Activities in Egypt; GH17-005, Conducting Public Health Research in China.

This publication corrects a notice that was published in the **Federal Register** on May 4, 2017, Volume 82, Number 85, pages 20894-20895. The meeting

announcement and matters for discussion should read as follows:

The meeting announced below concerns the Centers for Disease Control and Prevention (CDC) initial review of applications in response to Funding Opportunity Announcement (FOA) GH17-005, Conducting Public Health Research in China.

Time and Date: 9:00 a.m.–2:00 p.m., EDT, May 24, 2017

Matters for Discussion: The meeting will include the initial review, discussion, and evaluation of applications received in response to “Conducting Public Health Research in China”, GH17-005.

FOR FURTHER INFORMATION CONTACT:

Hylan Shoob, Scientific Review Officer, Center for Global Health (CGH) Science Office, CGH, CDC, 1600 Clifton Road, NE., Mailstop D-69, Atlanta, Georgia 30033, Telephone: (404) 639-4796, CGHERPO@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.

Elaine L. Baker,

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

[FR Doc. 2017-10334 Filed 5-19-17; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2017-N-1988]

Endocrinologic and Metabolic Drugs Advisory Committee; Notice of Meeting; Establishment of a Public Docket; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; establishment of a public docket; request for comments.

SUMMARY: The Food and Drug Administration (FDA or Agency) announces a forthcoming public advisory committee meeting of the Endocrinologic and Metabolic Drugs Advisory Committee. The general function of the committee is to provide advice and recommendations to the Agency on FDA’s regulatory issues. The meeting will be open to the public. FDA is establishing a docket for public comment on this document.

DATES: The meeting will be held on June 20, 2017, from 8 a.m. to 5 p.m. Comments received on or before June 6, 2017, will be provided to the committee. Comments received after that date will be taken into consideration by the Agency.

ADDRESSES: FDA White Oak Campus, 10903 New Hampshire Avenue, Building. 31 Conference Center, the Great Room (Rm. 1503), Silver Spring, MD 20993-0002. Answers to commonly asked questions including information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: <http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm>.

FDA is establishing a docket for public comment on this meeting. The docket number is FDA-2017-N-1988. All submissions received must include the Docket No. FDA-2017-N-1988 for “Endocrinologic and Metabolic Drugs Advisory Committee; Notice of Meeting; Establishment of a Public Docket; Request for Comments.” For detailed instructions on sending comments, see the “Public Participation” heading of the **SUPPLEMENTARY INFORMATION** section of this document. You may submit comments 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”).