

OFFICE OF GOVERNMENT ETHICS**Solicitation of Input From Stakeholders Regarding the U.S. Office of Government Ethics Strategic Plan (FY 2014–2017)**

AGENCY: Office of Government Ethics (OGE).

ACTION: Notice of Request for Public Comment.

SUMMARY: The U.S. Office of Government Ethics (OGE) is providing notice of request for public comment on its draft FY 2014–2017 Strategic Plan (Plan). The Plan describes OGE's priorities for the next four years. OGE will consider all comments received by the deadline. You may access the Plan at www.oge.gov, or you may obtain a copy of the Plan by sending an email request to OGEstrategicPlan@oge.gov.

DATES: All comments must be received on or before August 19, 2013.

ADDRESSES: You may submit comments by any of the following methods:

Email: OGEstrategicPlan@oge.gov.

Mail, Hand Delivery/Courier: U.S.

Office of Government Ethics, Suite 500, 1201 New York Avenue NW., Washington, DC 20005–3917, Attention: Nicole Stein, OGE Strategic Plan.

FOR FURTHER INFORMATION CONTACT:

Nicole Stein, Program Analyst, U.S. Office of Government Ethics, Suite 500, 1201 New York Avenue NW., Washington, DC 20005–3917; Telephone: 202–482–9255; TTY: 800–877–8339; Email: nicole.stein@oge.gov.

Dated: July 19, 2013.

Walter M. Shaub, Jr.

Director, U.S. Office of Government Ethics.

[FR Doc. 2013–17908 Filed 7–24–13; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES**Centers for Disease Control and Prevention****Advisory Board on Radiation and Worker Health (ABRWH or 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, announces the following meeting of the aforementioned committee:

Time and Date: 11:00 a.m.–3:00 p.m., September 5, 2013.

Place: Audio Conference Call via FTS Conferencing. The USA toll-free, dial-in

number is 1–866–659–0537 and the pass code is 9933701.

Status: Open to the public, but without a verbal public comment period. Written comment should be provided to the contact person below in advance of the meeting.

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, which 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 the CDC. NIOSH implements this responsibility for CDC. The charter was issued on August 3, 2001, renewed at appropriate intervals, most recently, August 3, 2011, and will expire on August 3, 2013.

Purpose: This 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, advising 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.

Matters To Be Discussed: The agenda for the conference call includes: Subcommittee and Work Group Updates; SEC Petition Evaluations Update for the October 2013 Advisory Board Meeting; Plans for the October 2013 Advisory Board Meeting; and Advisory Board Correspondence.

The agenda is subject to change as priorities dictate. Because there is not a public comment period, written

comments may be submitted. Any written comments received will be included in the official record of the meeting and should be submitted to the contact person below in advance of the meeting.

Contact Person for More Information: Theodore M. Katz, M.P.A., Designated Federal Official, NIOSH, CDC, 1600 Clifton Rd. NE., Mailstop: E–20, Atlanta, GA 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. 2013–17867 Filed 7–24–13; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES**Centers for Disease Control and Prevention****Clinical Laboratory Improvement Advisory Committee (CLIAC)**

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 of the aforementioned committee:

Times and Dates:

8:30 a.m.–5:00 p.m., August 21, 2013

8:30 a.m.–12:00 p.m., August 22, 2013

Place: CDC, 1600 Clifton Road NE., Tom Harkin Global Communications Center, Building 19, Room 232, Auditorium B, Atlanta, Georgia 30333.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 100 people.

Purpose: This Committee is charged with providing scientific and technical advice and guidance to the Secretary of Health and Human Services; the Assistant Secretary for Health; the Director, Centers for Disease Control and Prevention; the Commissioner, Food and Drug Administration; and the Administrator, Centers for Medicare and Medicaid Services. The advice and guidance pertain to general issues related to improvement in clinical laboratory quality and laboratory

medicine practice and specific questions related to possible revision of the CLIA standards. Examples include providing guidance on studies designed to improve safety, effectiveness, efficiency, timeliness, equity, and patient-centeredness of laboratory services; revisions to the standards under which clinical laboratories are regulated; the impact of proposed revisions to the standards on medical and laboratory practice; and the modification of the standards and provision of non-regulatory guidelines to accommodate technological advances, such as new test methods and the electronic transmission of laboratory information.

Matters To Be Discussed: The agenda will include agency updates from the CDC, the Centers for Medicare & Medicaid Services (CMS), and the Food and Drug Administration (FDA). Presentations and discussions will include improving laboratory quality in diverse settings, to include sites that perform waived testing as well as laboratories implementing telehealth initiatives such as digital pathology. Advancing laboratory interoperability in health information technology will also be discussed.

Agenda items are subject to change as priorities dictate.

Webcast: The meeting will also be Webcast. Persons interested in attending the in-person meeting or viewing the Webcast can access information about doing so at this URL: <http://www.cdc.gov/cliac/default.aspx>

Online Registration Required: All in-person CLIAC attendees are required to register for the meeting online at least 5 business days in advance for U.S. citizens and at least 10 business days in advance for international registrants. Register at <http://www.cdc.gov/cliac/default.aspx> by scrolling down and clicking the appropriate link under "Meeting Registration" (either U.S. Citizen Registration or Non-U.S. Citizen Registration) and completing all forms according to the instructions given. Please complete all the required fields before submitting your registration and submit no later than August 14, 2013 for U.S. registrants and August 7, 2013 for international registrants.

Providing Oral or Written Comments: It is the policy of CLIAC to accept written public comments and provide a brief period for oral public comments whenever possible. **Oral Comments:** In general, each individual or group requesting to make an oral presentation will be limited to a total time of five minutes (unless otherwise indicated). Speakers must also submit their comments in writing for inclusion in the

meeting's Summary Report. To assure adequate time is scheduled for public comments, individuals or groups planning to make an oral presentation should, when possible, notify the contact person below at least one week prior to the meeting date. **Written Comments:** For individuals or groups unable to attend the meeting, CLIAC accepts written comments until the date of the meeting (unless otherwise stated). However, it is requested that comments be submitted at least one week prior to the meeting date so that the comments may be made available to the Committee for their consideration and public distribution. Written comments, one hard copy with original signature, should be provided to the contact person below. Written comments will be included in the meeting's Summary Report.

Availability of Meeting Materials: To support the green initiatives of the federal government, the CLIAC meeting materials will be made available to the Committee and the public in electronic format (PDF) on the internet instead of by printed copy. Check the CLIAC Web site on the day of the meeting for materials. Note: If using a mobile device to access the materials, please verify the device's browser is able to download the files from the CDC's Web site before the meeting. Alternatively, the files can be downloaded to a computer and then emailed to the portable device. An internet connection, power source and limited hard copies may be available at the meeting location, but cannot be guaranteed. http://www.cdc.gov/cliac/cliac_meeting_all_documents.aspx

Contact Person for Additional Information: Nancy Anderson, Chief, Laboratory Practice Standards Branch, Division of Laboratory Science and Standards, Laboratory Science, Policy and Practice Program Office, Office of Surveillance, Epidemiology and Laboratory Services, CDC, 1600 Clifton Road NE., Mailstop F-11, Atlanta, Georgia 30329-4018; telephone (404) 498-2741; fax (404) 498-2210; or via email at NAnderson@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 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. 2013-17868 Filed 7-24-13; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2012-N-0977]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Regulations Restricting the Sale and Distribution of Cigarettes and Smokeless Tobacco To Protect Children and Adolescents

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Regulations Restricting the Sale and Distribution of Cigarettes and Smokeless Tobacco to Protect Children and Adolescents" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Daniel Gittleston, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50-400B, Rockville, MD 20850, 301-796-5156, Daniel.Gittleston@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: On April 10, 2013, the Agency submitted a proposed collection of information entitled "Regulations Restricting the Sale and Distribution of Cigarettes and Smokeless Tobacco to Protect Children and Adolescents" to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910-0312. The approval expires on July 31, 2016. A copy of the supporting statement for this information collection is available on the Internet at <http://www.reginfo.gov/public/do/PRAMain>.