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]

BILLING CODE 6345–03–P

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]

BILLING CODE 4163–18–P

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