staffing, and administrative support services for the Advisory Committee. The charter was reissued on May 12, 2015, and will expire on May 12, 2017.

Matters for Discussion: The Advisory Committee will continue its deliberations from the meeting on June 4, 2015, addressing the need for research on developmental or health effects in children. The agenda will include presentations on children's health research that has been conducted related to exposures from the 9/11 terrorist attacks. Also, a panel of children's researchers will discuss children's research issues with the Advisory Committee.

The agenda is subject to change as priorities dictate.

Public Comment Sign-up and Submissions to the Docket: To sign up to provide public comments or to submit comments to the docket, send information to the NIOSH Docket Office by one of the following means:

Mail: NIOSH Docket Office, Robert A. Taft Laboratories, C–34, 1090 Tusculum Avenue, Cincinnati, Ohio 45226.

Email: nioshdocket@cdc.gov.

Telephone: (513) 533–8611.

In the event an individual cannot attend, written comments may be submitted. The comments should be limited to two pages and submitted through *http://www.regulations.gov* by November 27, 2015. Efforts will be made to provide the two-page written comments received by the deadline below to the committee members before the meeting. Comments in excess of two pages will be made publicly available at *http://www.regulations.gov.* To view background information and previous submissions go to NIOSH docket *http://www.cdc.gov/niosh/docket/archive/docket248.html* , *http://www.cdc.gov/niosh/docket/archive/docket248–A.html,* and *http://www.cdc.gov/niosh/docket/archive/docket248–B.html.*

Policy on Redaction of Committee Meeting Transcripts (Public Comment): Transcripts will be prepared and posted to *http://www.regulations.gov* within 60 days after the meeting. If a person making a comment gives his or her name, no attempt will be made to redact that name. NIOSH will take reasonable steps to ensure that individuals making public comments are aware of the fact that their comments (including their name, if provided) will appear in a transcript of the meeting posted on a public Web site. Such reasonable steps include a statement read at the start of the meeting stating that transcripts will be posted and names of speakers will not be redacted. If individuals in making a statement reveal personal information (*e.g.,* medical information) about themselves, that information will not usually be redacted. The CDC Freedom of Information Act coordinator will, however, review such revelations in accordance with the Freedom of Information Act and, if deemed appropriate, will redact such information. Disclosures of information concerning third party medical information will be redacted.

Contact Person for More Information: Paul J. Middendorf, Ph.D., Designated Federal Officer, NIOSH, CDC, 2400 Century Parkway NE., Mail Stop E–20, Atlanta, GA 30345, telephone 1 (888) 982–4748; email: *wtc-stac@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.

Catherine Ramadei,

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

[FR Doc. 2015–26759 Filed 10–20–15; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Meeting: 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), and pursuant to the requirements of 42 CFR 83.15(a), the Centers for Disease Control and Prevention (CDC), announces the following meeting of the aforementioned committee:

Times and Dates: 8:15 a.m.–5 p.m., Pacific Time, November 18, 2015. 8:15 a.m.–4:30 p.m., Pacific Time, November 19, 2015.

Public Comment Time and Date: 5 p.m.–6 p.m.*, Pacific Time, November 18, 2015.

* Please note that the public comment period ends at the time indicated above or following the last call for comments, whichever is earlier. Members of the public who wish to provide public comments should plan to attend the public comment session at the start time listed.

Place: Waterfront Hotel, 10 Washington Street, Oakland, California 94607, Phone: 510–379–2652; Fax: 510–832–6228. Audio Conference Call via FTS Conferencing. The USA toll-free, dial-in number is 1–866–659–0537 with a pass code of 9933701. Live Meeting CONNECTION: *https://www.livemeeting.com/cc/cdc/join?id* GS59T7&role=attend&pw=ABRWH; Meeting ID: GS59T7; Entry Code: ABRWH.

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

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, and will expire on August 3, 2017.

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 For Discussion: The agenda for the Advisory Board meeting

includes: NIOSH Program Update; Department of Labor Program Update; Department of Energy Program Update; Report by the Dose Reconstruction Review Methods Work Group; SEC Petitions Update; Site Profile reviews for: Dow Chemical Co. (Madison, Illinois), and General Steel Industries (Granite City, Illinois); SEC petitions for: Battelle Laboratories, King Avenue (1956–1970; Columbus, Ohio), Lawrence Livermore National Laboratory (1974–1995; Livermore, California), Blockson Chemical Co. (1960–1991; Joliet, Illinois), Rocky Flats Plant (1984–1989; Golden, Colorado), Idaho National Laboratory (1949–1970; Scoville, Idaho), and Kansas City Plant (1949–1993; Kansas City, Missouri); and Board Work Sessions.

The agenda is subject to change as priorities dictate.

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

Policy on Redaction of Board Meeting Transcripts (Public Comment): (1) If a person making a comment gives his or her personal information, no attempt will be made to redact the name; however, NIOSH will redact other personally identifiable information, such as contact information, social security numbers, case numbers, etc., of the commenter.

(2) If an individual in making a statement reveals personal information (*e.g.,* medical or employment information) about themselves that information will not usually be redacted. The NIOSH Freedom of Information Act (FOIA) coordinator will, however, review such revelations in accordance with the Federal Advisory Committee Act and if deemed appropriate, will redact such information.

(3) If a commenter reveals personal information concerning a living third party, that information will be reviewed by the NIOSH FOIA coordinator, and upon determination, if deemed appropriated, such information will be redacted, unless the disclosure is made by the third party's authorized representative under the Energy Employees Occupational Illness Compensation Program Act (EEOICPA) program.

(4) In general, information concerning a deceased third party may be disclosed; however, such information will be redacted if (a) the disclosure is made by an individual other than the survivor claimant, a parent, spouse, or child, or the authorized representative of the deceased third party; (b) if it is unclear whether the third party is living or deceased; or (c) the information is unrelated or irrelevant to the purpose of the disclosure.

The Board will take reasonable steps to ensure that individuals making public comment are aware of the fact that their comments (including their name, if provided) will appear in a transcript of the meeting posted on a public Web site. Such reasonable steps include: (a) A statement read at the start of each public comment period stating that transcripts will be posted and names of speakers will not be redacted; (b) A printed copy of the statement mentioned in (a) above will be displayed on the table where individuals sign up to make public comments; (c) A statement such as outlined in (a) above will also appear with the agenda for a Board Meeting when it is posted on the NIOSH Web site; (d) A statement such as in (a) above will appear in the **Federal Register** Notice that announces Board and Subcommittee meetings.

Contact Person For More Information: Theodore Katz, Designated Federal Officer, NIOSH, CDC, 1600 Clifton Road NE., MS E–20, Atlanta, Georgia 30333, telephone: (513) 533–6800, toll free: 1–800–CDC–INFO, email: *dcas@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.

Catherine Ramadei,

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

[FR Doc. 2015–26757 Filed 10–20–15; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Meeting: Clinical Laboratory Improvement Advisory Committee

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 committee meeting.

Times and Dates

8:30 a.m.–5 p.m.., November 18, 2015.

8:30 a.m.–12 p.m., November 19, 2015.

Place: CDC, 2500 Century Center Boulevard, Rooms 1200/1201, Atlanta, Georgia 30345.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 100 people. This meeting will also be webcast, please see information below.

Purpose: This Committee is charged with providing scientific and technical advice and guidance to the Secretary of Health and Human Services (HHS); the Assistant Secretary for Health; the Director, Centers for Disease Control and Prevention; the Commissioner, Food and Drug Administration (FDA); and the Administrator, Centers for Medicare and Medicaid Services (CMS). 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 Clinical Laboratory Improvement Amendment (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 for Discussion: The agenda will include agency updates from CDC, CMS, and FDA. Presentations and discussions will include laboratory information exchange (interoperability); noninvasive prenatal testing; CLIA waiver guidance; the Institute of Medicine (IOM) report ''Improving Diagnosis in Health Care;'' and FDA guidance for laboratory developed tests.

Agenda items are subject to change as priorities dictate.

Webcast: The meeting will also be webcast. Persons interested in viewing the webcast can access information at: *http://cdclabtraining.adobeconnect.com/novcliac/*.

Online Registration Required: All people attending the CLIAC meeting in-person 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