exclusions portion of the Performance Information functional area, electronic records of past exclusions are maintained permanently in the archive list for historical reference. Federal agencies reporting exclusion information in SAM should follow their agency's guidance and policies for disposition of paper records.

SYSTEM MANAGER AND ADDRESS:

Integrated Award Environment Program Manager, Office of Integrated Award Environment, Federal Acquisition Service, U.S. General Services Administration, 2200 Crystal Drive, Arlington, Virginia 22202.

SAFEGUARDS:

System records are safeguarded in accordance with the requirements of the Privacy Act, the Computer Security Act, and the SAM System Security Plan. System roles are assigned with specific permissions to allow or prevent accessing certain information. Technical, administrative, and personnel security measures are implemented to ensure confidentiality and integrity of the system data that is stored, processed, and transmitted, including password protection and other appropriate security measures.

NOTIFICATION PROCEDURE:

For the Entity Management functional area, individuals know that SAM contains a record on them because they created the record. For the exclusions portion of the Performance Management functional area, individuals receive prior notification that their names will be contained in SAM from the Federal agency that takes the action to exclude them from Federal procurement and non-procurement programs. An individual may retrieve exclusion records by accessing the SAM public portal, which displays publicly available information only. Individuals may also contact the system program manager to inquire about any records about the individual.

RECORD ACCESS PROCEDURES:

Since individuals create the entity registration record in SAM and can delete or amend the record, there should not be any questions about that entry. However, individuals can contact the system manager with questions about the operation of the Entity Management functional area. Requests from individuals to determine the specifics of an exclusion record included in SAM should be addressed to the Federal agency POC identified in the exclusion record.

CONTESTING RECORD PROCEDURES:

Individuals or entities registered in SAM can edit their own registration record information. To contest the content of an exclusion record, individuals should contact the Federal agency point of contact identified in the exclusion record. For GSA provided exclusion records, procedures for contesting the content of a record and appeal procedures can be found at 41 CFR 105–64.

RECORD SOURCE CATEGORIES:

Entity records are created by the person or entity wishing to do business with the government. Exclusion records are created by Federal agency suspension and debarment personnel.

[FR Doc. 2013–03743 Filed 2–15–13; 8:45 am]

BILLING CODE 6820–34–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), 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:

Board Public Meeting Times and Dates (All times are Eastern Time): 9:45 a.m.–6:00 p.m., March 12, 2013.

Public Comment Times and Dates (All times are Eastern Time): 6:00 p.m.–7:00 p.m.,* March 12, 2012.

**Please note that the public comment period may end before the times indicated, following the last call for comments. Members of the public who wish to provide public comments should plan to attend public comment sessions at the start times listed.*

Place: Augusta Marriott Hotel, Two Tenth Street, Augusta, GA 30901; Phone: 706–722–8900; Fax: 706–724–0044. Audio Conference Call via FTS Conferencing. The USA toll-free, dial-in number is 1–866–659–0537 with a pass code of 9933701.

Status: Open to the public, limited only by the space available. The meeting space accommodates approximately 150 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, 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, 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.

Matters To Be Discussed: The agenda for the Advisory Board meeting includes: NIOSH Program Update; Department of Labor Program Update; Department of Energy Program Update; SEC petitions for: Brookhaven National Laboratory (1994–2007), Baker Brothers (Toledo, OH; 1945–1996); Procedures Review Subcommittee Report; SEC Issues Work Group Report on ''Sufficient Accuracy''; Savannah River Site Work Group Update; SEC Petitions Update; 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 in accordance with the redaction policy provided below. Any written comments received will be provided at the meeting and should be submitted to the contact person below well in advance of the meeting.

Policy on Redaction of Board Meeting Transcripts (Public Comment): (1) If a person making a comment gives his or her name, no attempt will be made to redact that name. (2) NIOSH 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. (3) If an individual in making a statement reveals personal information (e.g., medical 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 FOIA and the Federal Advisory Committee Act and if deemed appropriate, will redact such information. (4) All disclosures of information concerning third parties will be redacted. (5) If it comes to the attention of the DFO that an individual wishes to share information with the Board but objects to doing so in a public forum, the DFO will work with that individual, in accordance with the Federal Advisory Committee Act, to find a way that the Board can hear such comments.

Contact Person for More Information: Theodore Katz, DFO, NIOSH, CDC, 1600 Clifton Road, MS E–20, Atlanta, GA 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.

Elaine L. Baker,

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

[FR Doc. 2013–03612 Filed 2–15–13; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, Office of Infectious Diseases (BSC, OID)

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:

Time and Date: 3:00–4:00 p.m. Eastern Time, March 14, 2013.

Place: Teleconference.

Status: The meeting is open to the public; the toll free dial in number is 1–877–951–7311 with a passcode of 6420598.

Purpose: The BSC, OID, provides advice and guidance to the Secretary, Department of Health and Human Services; the Director, CDC; the Director, OID; and the Directors of the National Center for Immunization and Respiratory Diseases, the National Center for Emerging and Zoonotic Infectious Diseases, and the National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention, CDC, in the following areas: strategies, goals, and priorities for programs; research within the national centers; and overall strategic direction and focus of OID and the national centers.

Matters To Be Discussed: The purpose of the meeting is to discuss the potential for forming an infectious disease laboratory working group under the BSC, OID.

The agenda and any supplemental material will be available at *www.cdc.gov/oid/BSC.html* after March 1.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Robin Moseley, M.A.T., Designated Federal Officer, OID, CDC, 1600 Clifton Road NE., Mailstop D10, Atlanta, Georgia 30333, Telephone: (404) 639–4461.

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. 2013–03610 Filed 2–15–13; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Institute for Occupational Safety and Health (BSC, 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 of the aforementioned committee:

Time and Dates: 8:30 a.m.–3:15 p.m., March 21, 2013.

Place: Patriots Plaza I, 395 E Street SW., Room 9200, Washington, DC 20201.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 50 people. If you wish to attend in person, please contact NIOSH at (202) 245–0625 or (202) 245–0626 for information on building access. Teleconference is available toll-free; please dial (877) 328–2816, Participant Pass Code 6558291.

Purpose: The Secretary, the Assistant Secretary for Health, and by delegation the Director, Centers for Disease Control and Prevention, are authorized under Sections 301 and 308 of the Public Health Service Act to conduct directly or by grants or contracts, research, experiments, and demonstrations relating to occupational safety and health and to mine health. The Board of Scientific Counselors shall provide guidance to the Director, National Institute for Occupational Safety and Health on research and prevention programs. Specifically, the Board shall provide guidance on the Institute's research activities related to developing and evaluating hypotheses, systematically documenting findings and disseminating results. The Board shall evaluate the degree to which the activities of the National Institute for Occupational Safety and Health: (1) Conform to appropriate scientific standards, (2) address current, relevant needs, and (3) produce intended results.

Matters To Be Discussed: NIOSH Director Update; Implementation of the National Academies Program Recommendations for Construction Safety and Health, Respiratory Disease Studies, and Traumatic Injury Prevention, Nanotechnology Research Strategic Plan, Influenza Research, Agriculture, Forestry, and Fishing Sector Update.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Roger Rosa, Ph.D., Designated Federal Officer, BSC, NIOSH, CDC, 395 E Street SW., Suite 9200, Patriots Plaza Building, Washington, DC 20201, telephone (202) 245–0655, fax (202) 245–0664.

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 CDC 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. 2013–03737 Filed 2–15–13; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–0115]

Agency Information Collection Activities; Proposed Collection; Comment Request; Manufactured Food Regulatory Program Standards

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of