every effort to ensure that the membership of HHS Federal advisory committees is fairly balanced in terms of points of view represented and the committee’s function. Every effort is made that a broad representation of geographic areas, gender, ethnic and minority groups, and the disabled are given consideration for membership on HHS Federal advisory committees. Appointment to this committee shall be made without discrimination on the basis of age, race, ethnicity, gender, sexual orientation, disability, and cultural, religious, or socioeconomic status.

The Standards of Ethical Conduct for Employees of the Executive Branch are applicable to individuals who are appointed as public members of Federal advisory committees. Individuals appointed to serve as public members of Federal advisory committees are classified as special Government employees (SGEs). SGEs are Government employees for purposes of the conflict of interest laws. Therefore, individuals appointed to serve as public members of NVAC are subject to an ethics review. The ethics review is conducted to determine if the individual has any interests and/or activities in the private sector that may conflict with performance of their official duties as a member of the Committee. Individuals appointed to serve as public members of the Committee will be required to disclose information regarding financial holdings, consultancies, and research grants and/or contracts.

Bruce Gellin,
Deputy Assistant Secretary for Health,
Director, National Vaccine Program Office, Executive Secretary, National Vaccine Advisory Committee.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Board on Radiation and Worker Health (ABRW 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):**
  - 8:15 a.m.–5 p.m., December 7, 2011.
  - 8:15 a.m.–4 p.m., December 8, 2011.

**Public Comment Times and Dates (All times are Eastern Time):**
- 5 p.m.–6:30 p.m.,* December 7, 2011.

*Please note that the public comment periods 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:** Tampa Marriott Westshore, 1001 N. Westshore Blvd., Tampa, Florida 33607; Phone: (800) 564–3489; Fax: (813) 289–5464. 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 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, 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 Advisory Board meeting includes: NIOSH Program Update; Department of Labor (DOL) Program Update; Department of Energy (DOE) Program Update; Mound Plant Work Group Update; Pinellas Plant Site Profile Update; SEC petitions for: Weldon Spring, Hooker Electrochemical, Linde Ceramics Plant, Feed Materials Production Center (Fernald, Ohio), General Steel Industries; and Savannah River Site; SEC Petition Status Updates; Subcommittee and Work Group Reports; 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 to be posted on a public Web site. Reasonable steps include: (a) A person making a comment gives his or her name, no attempt will be made to redact that name; (b) 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 FOIA coordinator will, however, review such revelations in accordance with the Freedom of Information Act (5 U.S.C. 552).
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 National HIV Behavioral Surveillance For Young Men Who Have Sex With Men, Funding Opportunity Announcement (FOA), PS11–0010201SUPP12, initial review.

According 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 aforementioned meeting:

Time and Date

1 p.m.–5 p.m., January 12, 2012 (Closed).

Place: Teleconference.

Status: The meeting will be closed to the public in accordance with provisions set forth in Section 552b(c)(4) and (6), Title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92–463.

Matters To Be Discussed: The meeting will include the initial review, discussion, and evaluation of applications received in response to “National HIV Behavioral Surveillance For Young Men Who Have Sex With Men, FOA PS11–0010201SUPP12.”

Contact Person for More Information:

Amy Yang, Ph.D., Scientific Review Officer, CDC, 1600 Clifton Road, NE., Mailstop E609, Atlanta, Georgia 30333, Telephone: (404) 718–8836.

The Director, Management Analysis and Services Office, has 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.

Dated: November 9, 2011.

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

[FR Doc. 2011–29880 Filed 11–17–11; 8:45 am]
BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare & Medicaid Services (CMS) is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency’s functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

1. Type of Information Collection Request: New collection (request for new OMB control number); Title of Information Collection: Nursing Home Quality Improvement Questionnaire; Use: The information obtained via the Nursing Home Quality Improvement Questionnaire will be utilized by CMS staff in the Survey & Certification Group, Division of Nursing Homes, to identify areas for quality assurance and performance improvement (QAPI) technical assistance (TA) that will be useful to nursing facilities as they prepare to meet the new QAPI regulation that was mandated as part of the Affordable Care Act. Specifically, the information collected through the use of the questionnaire will be used to establish a baseline of QAPI practices in nursing homes, gather information on the challenges and barriers to implementing effective QAPI programs, assess the development of QAPI systems, determine what types of TA to make available to nursing homes, and assess the potential impact of TA in advancing QAPI in nursing homes; Form Number: CMS–10366 (OCN 0938–New); Frequency: Once; Affected Public: Private sector (business or other for-profits and not-for-profit institutions) and State, Local or Tribal Governments; Number of Respondents: 4,200; Total Annual Responses: 4,200; Total Annual Hours: 1,386. (For policy questions regarding this collection contact Debra Lyons at (410) 786–6780. For all other issues call (410) 786–1326.) To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS’ Web Site address at http://www.cms.hhs.gov/PaperworkReductionActof1995, or Email your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@cms.hhs.gov, or call the Reports Clearance Office on (410) 786–1326.

In commenting on the proposed information collections please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in one of the following ways by January 17, 2012:

1. Electronically. You may submit your comments electronically to http://www.regulations.gov. Follow the instructions for “Comment or Submission” or “More Search Options” to find the information collection document(s) accepting comments.

2. By regular mail. You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs,