

safety and health education, ergonomics, epidemiology, biostatistics, and psychology. Federal employees will not be considered for membership. Members may be invited to serve for up to four-year terms. Selection of members is based on candidates' qualifications to contribute to the accomplishment of the board's objectives <http://www.cdc.gov/niosh/BSC/default.html>.

DATES: Nominations for membership on the BSC must be received no later than August 1, 2018. Packages received after this time will not be considered for the current membership cycle.

ADDRESSES: All nominations should be mailed to NIOSH Docket 278, c/o Pauline Benjamin, Committee Management Specialist, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, 1600 Clifton Rd. NE, MS: E-20, Atlanta, Georgia 30329, or emailed (recommended) to nioshdocket@cdc.gov.

FOR FURTHER INFORMATION CONTACT: Alberto Garcia, M.S., DFO, CDC/NIOSH, 1090 Tusculum Ave. MS R-5, Cincinnati, OH 45226, telephone (513) 841-4596; agarcia1@cdc.gov.

SUPPLEMENTARY INFORMATION: The U.S. Department of Health and Human Services policy stipulates that committee membership be balanced in terms of points of view represented, and the committee's function. Appointments shall be made without discrimination on the basis of age, race, ethnicity, gender, sexual orientation, gender identity, HIV status, disability, and cultural, religious, or socioeconomic status. Nominees must be U.S. citizens, and cannot be full-time employees of the U.S. Government. Current participation on federal workgroups or prior experience serving on a federal advisory committee does not disqualify a candidate; however, HHS policy is to avoid excessive individual service on advisory committees and multiple committee memberships. Committee members are Special Government Employees (SGEs), requiring the filing of financial disclosure reports at the beginning and annually during their terms. CDC reviews potential candidates for NIOSH BSC membership each year, and provides a slate of nominees for consideration to the Secretary of HHS for final selection. HHS notifies selected candidates of their appointment near the start of the term in January 2019, or as soon as the HHS selection process is completed. Note that the need for expertise varies from year to year and a candidate who is not selected in one year may be reconsidered in a subsequent year. SGE Nominees must be

U.S. citizens, and cannot be full-time employees of the U.S. Government. Candidates should submit the following items:

- Current curriculum vitae, including complete contact information (telephone numbers, mailing address, email address).
- At least one letter of recommendation from person(s) not employed by the U.S. Department of Health and Human Services. (Candidates may submit letter(s) from current HHS employees if they wish, but at least one letter must be submitted by a person not employed by an HHS agency (e.g., CDC, NIH, FDA, etc.).

Nominations may be submitted by the candidate him- or herself, or by the person/organization recommending the candidate. 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 CDC and the Agency for Toxic Substances and Disease Registry.

Elaine Baker,

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

[FR Doc. 2018-12150 Filed 6-5-18; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Board on Radiation and Worker Health (ABRWH or the Advisory Board), Subcommittee on Dose Reconstruction Review (SDRR), National Institute for Occupational Safety and Health (NIOSH)

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of meeting.

SUMMARY: In accordance with the Federal Advisory Committee Act, the CDC announces the following meeting for the Subcommittee for Dose Reconstruction Reviews (SDRR) of the Advisory Board on Radiation and Worker Health (ABRWH). This meeting is open to the public, but without a public comment period. The public is welcome to submit written comments in advance of the meeting, to the contact person below. Written comments received in advance of the meeting will be included in the official record of the meeting. The public is also welcome to listen to the meeting by joining the teleconference at the USA toll-free, dial-

in number at 1-866-659-0537; the pass code is 9933701. The conference line has 150 ports for callers.

DATES: The meeting will be held on July 24, 2018, 10:30 a.m. to 4:30 p.m. EDT.

ADDRESSES: Audio Conference Call via FTS Conferencing. The USA toll-free dial-in number is 1-866-659-0537; the pass code is 9933701.

FOR FURTHER INFORMATION CONTACT:

Theodore Katz, MPA, Designated Federal Officer, NIOSH, CDC, 1600 Clifton Road, Mailstop E-20, Atlanta, Georgia 30329, Telephone (513)533-6800, Toll Free 1(800)CDC-INFO, Email ocas@cdc.gov.

SUPPLEMENTARY INFORMATION:

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 that 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 CDC. NIOSH implements this responsibility for CDC. The charter was issued on August 3, 2001, renewed at appropriate intervals, rechartered on February 12, 2018, pursuant to Executive Order 13708, and will terminate on September 30, 2019.

Purpose: The 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. SDRR was established to aid the Advisory Board in carrying out its duty to advise the Secretary, HHS, on dose reconstruction.

Matters To Be Considered: The agenda will include discussions on the following dose reconstruction program quality management and assurance activities: Dose reconstruction cases under review from Sets 19–24, including Iowa Ordinance Plant, Nevada Test Site, Los Alamos National Laboratory, Feeds Material Production Center (Fernald), Pantex Plant, Rocky Flats Plant, W.R. Grace, Hanford, Savannah River Site, Fernald, GE Evendale, Texas City Chemicals, Canoga Avenue Facility, De Soto Avenue Facility, Pacific Northwest National Laboratory, Amchitka Island Nuclear Explosion Site, Oak Ridge facilities, Paducah Gaseous Diffusion Plant, and potentially other Department of Energy and Atomic Weapons Employers facilities.

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 Baker,

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

[FR Doc. 2018–12149 Filed 6–5–18; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–18–0920; Docket No. CDC–2018–0053]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies the opportunity to comment on a proposed and/or continuing

information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled *Data Collection Through Web Based Surveys for Evaluating Act Against AIDS Social Marketing Campaign Phases Targeting Consumers* which includes web surveys to test campaign messaging.

DATES: CDC must receive written comments on or before August 6, 2018.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2018–0053 by any of the following methods:

- **Federal eRulemaking Portal:** *Regulations.gov.* Follow the instructions for submitting comments.
- **Mail:** Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS–D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to *Regulations.gov.*

Please note: Submit all comments through the Federal eRulemaking portal (*regulations.gov*) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Leroy A. Richardson, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS–D74, Atlanta, Georgia 30329; phone: 404–639–7570; Email: *omb@cdc.gov.*

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the

functions of the agency, including whether the information will have practical utility;

2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

3. Enhance the quality, utility, and clarity of the information to be collected; and

4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submissions of responses.

5. Assess information collection costs.

Proposed Project

Data Collection Through Web Based Surveys for Evaluating Act Against AIDS Social Marketing Campaign Phases Targeting Consumers—Extension—National Center for HIV/AIDS, Viral Hepatitis, STD and TB Prevention (NCHHSTP, Centers for Disease Control and Prevention (CDC).

Background and Brief Description

In response to the continued HIV epidemic in our country, CDC launched Act Against AIDS (AAA), a multifaceted communication campaign to reduce HIV incidence in the United States in 2009. CDC has released the campaign in phases, with some of the phases running concurrently. Each phase of the campaign uses mass media and direct-to-consumer channels to deliver messages. Some campaigns provide basic education and increase awareness of HIV/AIDS among the general public whereas others emphasize HIV prevention and testing among specific subgroups or communities at greatest risk of infection. CDC will also develop new messages to address changes in prevention science and subpopulations affected by HIV. The proposed study will assess the effectiveness of these social marketing messages aimed at increasing HIV/AIDS awareness, increasing prevention behaviors, and improving HIV testing rates among consumers.

This extension of an ongoing study will allow for continued evaluation of the effectiveness of AAA social marketing campaign through surveys with consumers. A total of 10,750 respondents were approved for the previously renewed generic ICR (0920–0920) and since the approval date, 4,305 respondents were surveyed under the GenIC, “Development of Messages for