

novel influenza viruses and biological agents that pose a threat to global public health. Through this international collaboration ASPR will capitalize on GMI's influential position as an important regional partner on disease surveillance efforts and public health emergency preparedness. This new cooperative agreement will help to further strengthen laboratory diagnostic capacity in Panama and other countries in the region. GMI's strong collaborative relationships with neighboring governments, as well as its training capabilities, and laboratory infrastructure will be critical for the viability of this partnership. In addition, this collaboration will support overall HHS efforts to continue building capacity abroad with the ultimate intent of detecting, stopping, slowing or otherwise limiting the threat or actual spread of bio-terrorism agents or a pandemic to the United States, thereby enhancing the health security of the American population.

ADDITIONAL INFORMATION: The agency program contact is Dr. Maria Julia Marinissen, who can be contacted at 202-205-4214 or Maria.Marinissen@hhs.gov.

Statutory Authority: Sections 301, 307, 1701 and 2811 of the Public Health Service Act, 42 U.S.C. 241, 2421, 300u, 300hh-10.

Dated: August 15, 2011.

Nicole Lurie,

Assistant Secretary for Preparedness and Response.

[FR Doc. 2011-21294 Filed 8-19-11; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Board on Radiation and Worker Health: Notice of Charter Renewal

This gives notice under the Federal Advisory Committee Act (Pub. L. 92-463) of October 6, 1972, that the Advisory Board on Radiation and Worker Health, Department of Health and Human Services, has been renewed for a 2-year period through August 3, 2013.

For information, contact Mr. Theodore Katz, Designated Federal Officer, Advisory Board on Radiation and Worker Health, Department of Health and Human Services, 1600 Clifton Road, M/S E20, Atlanta, Georgia 30341, telephone 404/498-2533, or fax 404/498-2570.

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.

Dated: August 12, 2011.

Elizabeth Millington,

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

[FR Doc. 2011-21413 Filed 8-19-11; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-11-11KA]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404-639-5960 and send comments to Catina Conner, CDC Acting Reports Clearance Officer, 1600 Clifton Road, MS-D74, Atlanta, GA 30333 or send an e-mail to omb@cdc.gov.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Written comments should be received within 60 days of this notice.

Proposed Project

Use of Evidence-Based Practices for Comprehensive Cancer Control—New—National Center for Chronic Disease

Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

There have been increasing calls in the fields of public health generally and cancer control specifically for the dissemination, adoption, and implementation of evidence-based practices (EBPs). EBPs are public health practices (interventions, programs, strategies, policies, procedures, processes, and/or activities) that have been tested or evaluated and shown to be effective. However, while the development, review, and compilation of EBPs has steadily increased over time, there is concern that the adoption and implementation of those practices, including among cancer control planners and practitioners, has not kept pace. Given the gap between the development of EBPs and their use, public health and cancer control organizations need to place greater emphasis on the promotion and dissemination of these practices among those who can use them to improve population health.

While efforts to promote cancer control EBPs have increased, questions remain whether these efforts will result in widespread adoption and implementation of EBPs in the context of comprehensive cancer control (CCC) in the states, Tribes, and U.S. Associated Pacific Island Jurisdictions and territories. National Comprehensive Cancer Control Program (NCCCCP) grantees may face a number of challenges to incorporating EBPs into CCC efforts in their jurisdictions. In order to address these barriers effectively and better promote the use of EBPs for cancer control, CDC would like to understand (1) how evidence-based approaches are currently being used to develop CCC plans; (2) how CCC programs identify EBPs; (3) what EBPs have been adopted by CCC programs; and (4) what challenges and unintended consequences have been encountered in their implementation.

CDC plans to conduct a new, one-time study to examine CCC planners' use of scientific and practice-based information to inform development of their CCC plans. Information collection will consist of two Web-based surveys involving key CCC stakeholders in the NCCCCP-funded states, Tribes, and U.S. Associated Pacific Island Jurisdictions and territories. Respondents for the first survey will be Directors of the 66 NCCCCP-funded programs, who will also have the opportunity to participate in a follow-up telephone call. Respondents for the second survey will be key

program partners/collaborators identified by each Program Director (1–2 partners per Director) as instrumental to the selection and implementation of cancer control EBPs. The survey results will help CDC enhance existing NCCCP efforts by identifying new strategies for promoting the use of evidence-based approaches to comprehensive cancer control. The surveys will also identify

technical assistance needs of NCCCP-funded awardees related to selection and implementation of EBPs, and will contribute to CDC's efforts to build the capacities of states, Tribes, and Pacific Island Jurisdictions and territories toward more effective efforts in cancer prevention and control. Finally, the results may lead to new insights and questions that can be addressed in

future studies. CDC's authorization to conduct the study is provided by Section 301 of the Public Health Service Act (42 U.S.C. 241).

OMB approval will be requested for one year. Participation in the study is voluntary. There are no costs to the respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of Respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total response burden (in hours)
CCC Program Directors	Survey Scheduling Script	66	1	15/60	17
	Program Directors Web Survey Questionnaire.	66	1	0.5	33
	Program Directors Telephone Interview Guide and Script.	66	1	20/60	22
CCC Program Partners	Program Partners Web Survey Questionnaire.	132	1	0.5	66
Total	138

Catina Conner,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. 2011–21400 Filed 8–19–11; 8:45 am]

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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 Member Conflict Review, Program Announcement (PA) 07–318, initial review.

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 aforementioned meeting:

Time and Date: 1 p.m.–3 p.m., November 9, 2011 (Closed).

Place: National Institute for Occupational Safety and Health (NIOSH), CDC, 1095 Willowdale Road, Morgantown, West Virginia 26506, *Telephone:* (304) 285–6143.

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 “Member Conflict Review, PA 07–318.”

Contact Person for More Information:

Bernadine Kuchinski, PhD, Scientific Review Officer, Office of Extramural Programs, NIOSH, CDC, Robert A. Taft Laboratories, 4676 Columbia Parkway, MS C–7, Cincinnati, Ohio 45226, *Telephone:* (513) 533–8511.

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.

Dated: August 12, 2011.

Elizabeth Millington,

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

[FR Doc. 2011–21420 Filed 8–19–11; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Safety and Occupational Health Study Section (SOHSS), 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 (CDC), announces the following meeting of the aforementioned committee:

Times and Dates:

8 a.m.–5 p.m., October 18, 2011 (Closed).

8 a.m.–5 p.m., October 19, 2011 (Closed).

Place: Embassy Suites, 1900 Diagonal Road, Alexandria, Virginia 22314, *Telephone:* (703) 684–5900, *Fax:* (703) 684–0653.

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.

Purpose: The Safety and Occupational Health Study Section will review, discuss, and evaluate grant application(s) received in response to the Institute's standard grants review and funding cycles pertaining to research issues in occupational safety and health, and allied areas.

It is the intent of NIOSH to support broad-based research endeavors in keeping with the Institute's program goals. This will lead to improved understanding and appreciation for the magnitude of the aggregate health burden associated with occupational injuries and illnesses, as well as to support more focused research projects, which will lead to improvements in the delivery of occupational safety and health services, and the prevention of work-related injury and illness. It is anticipated that research funded will promote these program goals.

Matters To Be Discussed: The meeting will convene to address matters related to the conduct of Study Section business and for the study section to consider safety and occupational health-related grant applications.

These portions of 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, Centers for Disease Control and Prevention, pursuant to Section 10(d) Pub. L. 92–463.