Leroy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-15-14ADD]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (a) 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; (b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (c) Enhance the quality, utility, and clarity of the information to be collected; (d) 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 submission of responses; and (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570 or send an email to *omb@cdc.gov*. Written comments and/or suggestions regarding the items contained in this notice should be directed to the Attention: CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

National Occupational Research Agenda (NORA) 2016 Decade Review— New—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The National Institute for Occupational Safety and Health (NIOSH) is responsible for conducting research and making recommendations to prevent worker injury and illness, as authorized in Section 20(a)(1) of the Occupational Safety and Health Act (29 U.S.C. 669). In 1995–6, NIOSH saw an opportunity to enhance its ability to accomplish its mission through partnerships that involved a broad national stakeholder base in occupational safety and health. With stakeholder input, NIOSH developed and launched a decade-long partnership program titled the National Occupational Research Agenda (NORA) in 1996. Participation in NORA includes stakeholders from universities, large and small businesses, professional societies, government agencies, and worker organizations. After an internal management review of the first decade of NORA, conducted in 2005, NIOSH launched the second decade of NORA (2006-2016) structured for even greater national impact. This information collection is a necessary part of a larger internal NIOSH management review of the second decade of NORA. The results of this review will inform NIOSH

decisions about how to structure a third decade of NORA (2016–2026) for maximum effectiveness and impact.

The second decade of NORA was based on a new sector structure to better move research to practice within workplaces. The work of the sectors is managed through a partnership structure of sector councils. Each council develops and maintains an agenda for the decade for its sector. The sector agendas become part of the national agenda for improvements in occupational safety and health through research and partnerships. Representing all stakeholders, the councils use an open process to set goals, develop strategies, encourage partnerships, and promote improved workplace practices.

NIOSH is requesting a 12-month OMB approval to administer a survey to NORA council members and leaders. The collection of information is necessary for NIOSH management to assess the efficiency and effectiveness of the NORA sector councils. The target population is all current and former members and leaders of each of the ten NORA Sector Councils. The web-based questionnaire requests information on satisfaction with the efficiency of the council and its processes, on impacts made in the sector during the second decade, and suggestions for improving the effectiveness and impact of NORA in the future. Without this data collection, NIOSH's internal management review of NORA would lack critical stakeholder input from its many non-Federal partners.

A 16-item questionnaire has been developed and will be sent to all 352 non-Federal NORA Sector council members or leaders. A pilot test of the questionnaire was conducted by asking eight NIOSH employees who are a leader of a NORA sector council to complete the questionnaire and provide feedback. Respondents to the pilot test estimated the questionnaire requires approximately 15 minutes to complete. The total estimated burden is 88 hours. There is no cost to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)
Council member or leader	Council Questionnaire	352	1	15/60

Leroy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2014–26132 Filed 11–3–14; 8:45 am] BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Committee on Breast Cancer in Young Women (ACBCYW)

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: 9:00 a.m.—5:30 p.m. EST, December 4, 2014; 8:00 a.m.—12:00 p.m. EST, December 5, 2014.

Place: CDC, 4770 Buford Highway, Chamblee Building 107 1B/1C, Atlanta, Georgia 30341.

This meeting is also accessible by teleconference and web access. Teleconference and web access login information is as follows:

Toll-Free Telephone: 1–800–621–3587,
Participant passcode: 9679129. There is also a toll number for anyone outside of the USA: TOLL NUMBER: 1–517–308–9263,
Participant passcode: 96791NET
CONFERENCE AND WEB URL: For
December 4, 2014: https://
www.mymeetings.com/nc/join/ Conference number: PW9240020, Audience passcode: 9679129

For December 5, 2014: https://www.mymeetings.com/nc/join/ Conference number: PW9240028, Audience passcode: 9679129.

Participants can join the event directly at: https://www.mymeetings.com/nc/join.php?i=PW9240020&p=9679129&t=c.

Status: Open to the public, limited only by the space and phone lines available.

Purpose: The committee provides advice and guidance to the Secretary, HHS; the Assistant Secretary for Health; and the Director, CDC, regarding the formative research, development, implementation and evaluation of evidence-based activities designed to prevent breast cancer (particularly among those at heightened risk) and promote the early detection and support of young women who develop the disease. The advice provided by the Committee will assist in ensuring scientific quality, timeliness, utility, and dissemination of credible appropriate messages and resource materials.

Matters for Discussion: The agenda will include discussions on the current and emerging topics related to breast cancer in young women. These may include public health communication, breast cancer in

young women digital and social media campaigns, and CDC updates. Topics will address efforts increase awareness around breast cancer risk, breast health, symptoms, diagnosis, and treatment of breast cancer in young women.

Agenda items are subject to change as priorities dictate.

Online Registration Required: In order to expedite the security clearance process required for entry into a Federal building, all ACBCYW attendees must register for the meeting online at least 7 days in advance at http://www.cdc.gov/cancer/breast/what_cdc is <a href="doi:no:

Contact Person for More Information: Temeika L. Fairley, Ph.D., Designated Federal Officer, National Center for Chronic Disease Prevention and Health Promotion, CDC, 5770 Buford Hwy. NE., Mailstop K52, Atlanta, Georgia 30341, Telephone (770) 488–4518, Fax (770) 488–4760. Email: acbcyw@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 Agency for Toxic Substances and Disease Registry.

Claudette Grant,

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

[FR Doc. 2014–26082 Filed 11–3–14; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

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

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: 10:00–11:00 a.m. EST, November 20, 2014.

Place: Teleconference.

Status: The meeting is open to the public; the toll free dial in number is 1–877–951–7311 with a pass code of 1588067.

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 for Discussion: Discussions will focus on current responses and national center priorities.

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.

Claudette Grant,

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

[FR Doc. 2014–26083 Filed 11–3–14; 8:45 am] BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS-10527]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. 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 information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates 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