

SUPPLEMENTARY INFORMATION:**A. Purpose**

The information collection activity will garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Administration's commitment to improving service delivery. By qualitative feedback we mean information that provides useful insights on perceptions and opinions, but are not statistical surveys that yield quantitative results that can be generalized to the population of study.

This feedback will provide insights into customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative and actionable communications between the Agency and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

Feedback collected under this generic clearance will provide useful information, but it will not yield data that can be generalized to the overall population. This type of generic clearance for qualitative information will not be used for quantitative information collections that are designed to yield reliably actionable results, such as monitoring trends over time or documenting program performance.

Such data uses require more rigorous designs that address: The target population to which generalizations will be made, the sampling frame, the sample design (including stratification and clustering), the precision requirements or power calculations that justify the proposed sample size, the expected response rate, methods for assessing potential non-response bias, the protocols for data collection, and any testing procedures that were or will be undertaken prior fielding the study.

Depending on the degree of influence the results are likely to have, such collections may still be eligible for submission for other generic mechanisms that are designed to yield quantitative results. The Digital Government Strategy released by the White House in May, 2012 drives agencies to have a more customer-centric focus. Because of this, GSA anticipates an increase in requests to use this generic clearance, as the plan states that: A customer-centric principle charges us to do several things: Conduct

research to understand the customer's business, needs and desires; "make content more broadly available and accessible and present it through multiple channels in a program-and device-agnostic way; make content more accurate and understandable by maintaining plain language and content freshness standards; and offer easy paths for feedback to ensure we continually improve service delivery.

The customer-centric principle holds true whether our customers are internal (e.g., the civilian and military federal workforce in both classified and unclassified environments) or external (e.g., individual citizens, businesses, research organizations, and state, local, and tribal governments)."

B. Annual Reporting Burden

Respondents: 500,000.

Responses per Respondent: 1.

Total Annual Responses: 500,000.

Hours per response: 60.446 minutes.

Total Burden hours: 32,970.72.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the GSA Regulatory Secretariat Division, by calling 202-501-4755 or emailing GSARegSec@gsa.gov. Please cite OMB Control No. 3090-0297, Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery, in all correspondence.

Beth Anne Killoran,
Deputy Chief Information Officer.

[FR Doc. 2022-05406 Filed 3-14-22; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES**Centers for Disease Control and Prevention****Notice of Award of a Single-Source Cooperative Agreement To Fund Burkina Faso Ministry of Health; Cancellation**

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

ACTION: Notice.

SUMMARY: The Centers for Disease Control and Prevention (CDC), located within the Department of Health and Human Services (HHS), announces the cancellation of an award of approximately \$450,000 for Year 1 of funding to the Burkina Faso Ministry of Health.

DATES: The notice of award was cancelled on March 10, 2022.

FOR FURTHER INFORMATION CONTACT: Trong Ao, Center for Global Health, Centers for Disease Control and Prevention, CDC Ghana Office, US Embassy, 24 Fourth Circular Road Cantonments, Accra Ghana, Telephone: 800-232-6348, E-Mail: tfa8@cdc.gov.

SUPPLEMENTARY INFORMATION: On March 8, 2022 CDC announced a single-source award will support the Burkina Faso Ministry of Health to implement HIV strategic information and laboratory strengthening activities in Burkina Faso (87 FR 12966). This award is cancelled in its entirety.

Terrance Perry,
Chief Grants Management Officer, Centers for Disease Control and Prevention.

[FR Doc. 2022-05565 Filed 3-11-22; 4:15 pm]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES**Centers for Disease Control and Prevention****Advisory Board on Radiation and Worker Health (ABRWH), Subcommittee for Dose Reconstruction Reviews (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 or the Advisory Board). 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.

DATES: The meeting will be held on April 20, 2022, 11:00 a.m. to 4:00 p.m., EDT. Written comments must be received on or before April 13, 2022.

ADDRESSES: You may submit comments by mail to: Sherri Diana, National Institute for Occupational Safety and Health, 1090 Tusculum Avenue, MS C-34, Cincinnati, Ohio 45226.

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

Written comments received in advance of the meeting will be included in the official record of the meeting. The public is also welcomed to listen to the meeting by joining the audio conference (information below). The audio conference line has 150 ports for callers.

FOR FURTHER INFORMATION CONTACT:

Rashaun Roberts, Ph.D., Designated Federal Officer, NIOSH, CDC, 1090 Tusculum Avenue, Mailstop C-24, Cincinnati, Ohio 45226, Telephone: (513) 533-6800, 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, and rechartered under Executive Order 13889 on March 22, 2020, and will terminate on March 22, 2022.

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 Set 30, possibly including cases involving the Hanford site and discussion of changes in SDRR selection criteria based on SC&A document, "Summary Dose Reconstruction Information," dated December 20, 2021. Agenda items are subject to change as priorities dictate. For additional information, please contact Toll Free 1 (800) CDC-INFO.

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, 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.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Center for Injury Prevention and Control, (BSC, NCIPC)

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 Board of Scientific Counselors, National Center for Injury Prevention and Control, (BSC, NCIPC). This is a virtual meeting and is open to the public, limited by the capacity of the

conference webinar which is 2,000 participants. Pre-registration is required. Time will be available for public comment.

DATES: The meeting will be held on April 11, 2022, from 10:00 a.m. to 4:30 p.m., EDT. The public may submit written comments from March 15, 2022 through April 18, 2022.

ADDRESSES: Zoom Virtual Meeting. If you would like to attend the virtual meeting, please pre-register by accessing the link at https://dceproductions.zoom.us/webinar/register/WN_Mi3IYbnQTOmuEKZBxScBWQ. Instructions to access the Zoom virtual meeting will be provided in the link following registration.

Meeting Information: There will be a public comment period from 3:45 p.m.–4:15 p.m., EDT. The public is encouraged to register for the BSC, NCIPC April 11, 2022, meeting public comment period by accessing the link provided: <https://www.surveymonkey.com/r/l3hdzb6>. Individuals wishing to pre-register for public comment must do so by Wednesday, April 6, 2022, at 5:00 p.m., EDT.

FOR FURTHER INFORMATION CONTACT:

Arlene Greenspan, DrPH, MPH, PT, Associate Director for Science, National Center for Injury Prevention and Control, Centers for Disease Control and Prevention, 4770 Buford Highway NE, Mailstop S106-9, Atlanta, Georgia 30341, Telephone: (770) 488-1279; Email address: ncipcbsc@cdc.gov.

SUPPLEMENTARY INFORMATION:

Purpose: The Board will: (1) Conduct, encourage, cooperate with, and assist other appropriate public health authorities, scientific institutions, and scientists in the conduct of research, investigations, experiments, demonstrations, and studies relating to the causes, diagnosis, treatment, control, and prevention of injury and violence, and other impairments; (2) assist States and their political subdivisions in preventing intentional and unintentional injuries and to promote health and well-being; and (3) conduct and assist in research and control activities related to injury. The Board of Scientific Counselors makes recommendations regarding policies, strategies, objectives, and priorities; and reviews progress toward injury prevention goals and provides evidence in injury prevention-related research and programs. In addition, the Board provides advice on the appropriate balance of intramural and extramural research, the structure, progress, and performance of intramural programs.