

September 30, 2025, consistent with E.O. 14109 (September 29, 2023).

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. The ABRWH Subcommittee on Procedure Reviews (SPR) is responsible for overseeing, tracking, and participating in the reviews of all procedures used in the dose reconstruction process by the NIOSH Division of Compensation Analysis and Support (DCAS) and its dose reconstruction contractor (Oak Ridge Associated Universities—ORAU).

Matters to be Considered: The meeting agenda will include discussions on the following: (1) Carry-over items from March 14, 2024 SPR meeting, including a. DCAS–PER–040 “Mallinckrodt TBD Revisions,” b. DCAS–PER–068 “Electro Metallurgical Co”, c. DCAS–PER–070 “Nuclear Metals Inc.”, d. DCAS–PER–072 “Seymour Specialty Wiring Co”, e. ORAUT–RPRT–0060 “Neutron Dose from Highly Enriched Uranium”, and f. DR template reviews—findings versus observations; (2) Newly issued SC&A reviews, including a. ORAUT–OTIB–0036 “Internal Dosimetry Coworker Data for Portsmouth Gaseous Diffusion Plant” b. ORAUT–OTIB–0040 “External Coworker Dosimetry Data for the Portsmouth Gaseous Diffusion Plant” c. ORAUT–OTIB–0093 “Conversion of Committed Effective Dose to Annual Organ Dose” and d. ORAUT–RPRT–0087 “Applications of Regression in External Dose Reconstruction”; (3) Preparation for August 2024 Full ABRWH Meeting: Review of technical guidance documents ready for full Board approval; (4) Newly Issued Guidance and Supplemental Topics. Agenda items are subject to change as priorities dictate. For additional information, please contact Toll Free 1(800) 232–4636.

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

[FR Doc. 2024–21497 Filed 9–19–24; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Meeting of the Advisory Board on Radiation and Worker Health, National Institute for Occupational Safety and Health

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 Centers for Disease Control and Prevention (CDC) announces the following meeting of the Advisory Board on Radiation and Worker Health (ABRWH or the Advisory Board). This is a virtual meeting. It 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 listed in the addresses section below. 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 teleconference (information below), limited only by the number of audio conference lines available (150).

DATES: The meeting will be held on October 9, 2024, from 11 a.m. to 1 p.m., EDT.

Written comments must be received on or before October 2, 2024.

ADDRESSES: You may submit comments by mail to: Rashaun Roberts, Ph.D., National Institute for Occupational Safety and Health, 1090 Tusculum Avenue, Mailstop C–24, 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.

FOR FURTHER INFORMATION CONTACT: Rashaun Roberts, Ph.D., Designated

Federal Officer, National Institute for Occupational Safety & Health, Centers for Disease Control and Prevention, 1090 Tusculum Avenue, Mailstop C–24, Cincinnati, Ohio 45226, Telephone (513) 533–6800, Toll Free 1(800) 232–4636, 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 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 rechartered under Executive Order 14109 (September 29, 2023) on March 22, 2024. Unless continued by the President the Board will terminate on September 30, 2025, consistent with E.O. 14109 of (September 29, 2023).

Purpose: This Advisory Board is charged with (a) providing advice to the Secretary, HHS, on the development of guidelines under E.O. 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 Considered: The agenda will include discussions on the following: Update on Cybersecurity Modernization Initiative; Work Group and Subcommittee reports; Update on

the status of SEC Petitions; and plans for the December 2024 Advisory Board Meeting. Agenda items are subject to change as priorities dictate. For additional information, please contact Toll Free 1(800) 232-4636.

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Kalwant Smagh,

Director, Office of Strategic Business Initiatives, 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

[30Day-24-24FA]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled “Human-Centered Design Effort on Bringing Guidelines to the Digital Age” to the Office of Management and Budget (OMB) for review and approval. CDC previously published a “Proposed Data Collection Submitted for Public Comment and Recommendations” notice on May 7, 2024 to obtain comments from the public and affected

agencies. CDC received two comments related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

(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.

Comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function. Direct written comments and/or suggestions regarding the items contained in this notice to the

Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street NW, Washington, DC 20503 or by fax to (202) 395-5806. Provide written comments within 30 days of notice publication.

Proposed Project

Human-Centered Design Effort on Bringing Guidelines to the Digital Age—Existing Collection in Use Without an OMB Control Number—Office of Public Health Data, Surveillance, and Technology (OPHDST), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Given the increased demand to improve clinical guideline development and implementation, a new approach that began with an initiative on Adapting Clinical Guidelines for the Digital Age has been expanded by Guidelines International Network North America to implement a future state of guideline development and implementation that leverages advancements in technology. To identify pain points in the process, there were discussions with individuals from multiple perspectives in guidelines development and implementation.

CDC requests approval for an Existing Collection in Use Without an OMB Control Number, for Human-Centered Design Effort on Bringing Guidelines to the Digital Age. Data from this project will be used to inform the structure of a human-centered design workshop where participants use the pain points identified from the semi-structured interviews as the starting point for exploring insights about guideline development and implementation.

The burden estimates include the time for respondents to be interviewed. The estimated annual burden for respondents 33 hours. There is no cost to respondents other than their time to participate.

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Clinicians	Clinician Conversation Guide	5	1	1
EHR Vendors	EHR Vendor Conversation Guide	2	1	1
Guideline Developers	Guideline Developer Conversation Guide	8	1	1
Informaticists	Informaticist Conversation Guide	4	1	1
Implementers	Implementer Conversation Guide	9	1	1
Insurers	Insurer Conversation Guide	1	1	1
Patient/Patient Advocate	Patient/Patient Advocate Conversation Guide	4	1	1