the same or similar items and other references (including contract numbers, points of contact with telephone numbers and other relevant information).

Postaward. For responses in the Contractor Performance Assessment Reporting System (CPARS).

• *FAR 42.1503(d).* Requires contractors be afforded up to 14 calendar days from the notification date that a past performance evaluation has been entered into CPARS to submit comments, rebutting statements, or additional information. Past performance information is relevant information regarding a contractor's actions under previously awarded contracts or orders, for future source selection purposes. Source selection officials may obtain past performance information from a variety of sources.

The contracting officer will use the information to support future source selection decisions.

C. Annual Burden

Respondents: 65,373.
Total Annual Responses: 83,262.
Total Burden Hours: 166,524.

D. Public Comment

A 60-day notice was published in the **Federal Register** at 86 FR 8913, on February 10, 2021. One respondent submitted comments; however, they did not change the estimate of the burden.

Comment: The commenter expressed there is a need for a revision to the practices of acquiring, analyzing, and utilizing past performance information for source selection in federal acquisitions. The commenter noted that ''there have emerged in recent years a number of issues that result in a misleading, incomplete, deceptive, or distorted assessment of an Offeror's Past Performance that is not allowing fair competition for contracts which evaluate this factor.'' The commenter suggested changes on a few areas by using examples.

Response: Some of the suggestions made by the commenter may require consideration via the rulemaking process, and other suggestions refer to practices by particular agencies. These suggestions are outside the scope of this information collection renewal. The commenter did not express an opinion on whether the collections of information are needed; whether the estimated number of burden hours is accurate; or ways to minimize the burden of the collection of information. Therefore, the estimate of the burden was not changed.

Obtaining Copies: 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. 9000–0142, Past Performance Information.

Janet Fry,

Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2021–08409 Filed 4–22–21; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Board on Radiation and Worker Health (ABRWH), 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 welcomed to listen to the meeting by joining the audio conference (information below). The audio conference line has 150 ports for callers.

DATES: The meeting will be held on June 16, 2021, from 10:30 a.m. to 2:30 p.m., EDT. Written comments must be received on or before June 9, 2021.

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.

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, 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 Advisory Board's charter was issued on August 3, 2001, renewed at appropriate intervals, rechartered 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 29, possibly including cases involving: Albuquerque Operations Office, Area IV of the Santa Susana Field Laboratory, Argonne National Laboratory-East, Argonne National Laboratory-West, Battelle Laboratories-King Avenue, Clarksville Modification Center, Feed Materials Production Center (FMPC), Fermi National Accelerator Laboratory, General Atomics, Hanford, Idaho National Laboratory, Lawrence Berkeley National Laboratory, Lawrence Livermore National Laboratory, Los Alamos National Laboratory, Mound Plant, Nevada Test Site, Oak Ridge Gaseous Diffusion Plant (K–25), Oak Ridge Institute for Science Education, Oak Ridge National Laboratory (X–10), Pacific Northwest National Laboratory, Paducah Gaseous Diffusion Plant, Pantex Plant, Portsmouth Gaseous Diffusion Plant, Rocky Flats Plant, Savannah River Site, and/or Y–12 Plant. Agenda items are subject to change as priorities dictate.

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.

[FR Doc. 2021–08428 Filed 4–22–21; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Mine Safety and Health Research Advisory Committee (MSHRAC)

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 Mine Safety and Health Research Advisory Committee (MSHRAC). This is a virtual meeting. It is open to the public, limited only by web conference lines (500 web conference lines are available).

DATES: The meeting will be held on June 21, 2021, from 10:00 a.m. to 2:30 p.m., EDT.

ADDRESSES: If you wish to attend the virtual meeting, please contact Ms. Berni Metzger by email at *metzger@cdc.gov* or by telephone at (412) 386–4541 at least 5 business days in advance of the meeting. She will provide you with the Zoom web conference access information.

FOR FURTHER INFORMATION CONTACT: George W. Luxbacher, Designated Federal Officer, MSHRAC, National Institute for Occupational Safety and Health (NIOSH), CDC, 2400 Century Parkway NE, Atlanta, GA 30345, Telephone: (404) 498–2808; Email: *gluxbacher@cdc.gov.*

SUPPLEMENTARY INFORMATION:

Purpose: This committee is charged with providing advice to the Secretary, Department of Health and Human Services; the Director, CDC; and the Director, NIOSH, on priorities in mine safety and health research, including grants and contracts for such research, 30 U.S.C. 812(b)(2), Section 102(b)(2).

Matters To Be Considered: The agenda will include discussions on NIOSH mining safety and health research capabilities, projects, and outcomes, including FY21 new mining projects; updates on MINER Act extramural research; and current intramural dust, diesel particulate matter (DPM) and silica research. The meeting will also include an update from the NIOSH Associate Director for Mining. Agenda items are subject to change as priorities dictate.

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.

[FR Doc. 2021–08429 Filed 4–22–21; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10500]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human 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 the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by June 22, 2021.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to *http://www.regulations.gov.* Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number: CMS–P–0015A, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.