DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Notice of Meetings

AGENCY: Agency for Healthcare Research and Quality (AHRQ), Health and Human Services (HHS).

ACTION: Notice of five AHRQ subcommittee meetings.

SUMMARY: The subcommittees listed below are part of AHRQ's Health Services Research Initial Review Group Committee. Grant applications are to be reviewed and discussed at these meetings. Each subcommittee meeting will be closed to the public.

DATES: See below for dates of meetings:

1. *Healthcare Safety and Quality Improvement Research (HSQR)*
   Date: October 14th, 2020
2. *Healthcare Effectiveness and Outcomes Research (HEOR)*
   Date: October 14–15, 2020
3. *Health Care Research and Training (HCRT)*
   Date: October 15–16, 2020
4. *Health System and Value Research (HSVR)*
   Date: October 15–16, 2020
5. *Healthcare Information Technology Research (HITR)*
   Date: October 22–23, 2020

ADDRESSES: Agency for Healthcare Research and Quality (Virtual Review), 5600 Fishers Lane, Rockville, Maryland 20857.

FOR FURTHER INFORMATION CONTACT: (To obtain a roster of members, agenda or minutes of the non-confidential portions of the meetings.)

Jenny Griffith, Committee Management Officer, Office of Extramural Research Education and Priority Populations, Agency for Healthcare Research and Quality (AHRQ), 5600 Fishers Lane, Rockville, Maryland 20857, Telephone (301) 27–1557.

SUPPLEMENTARY INFORMATION: In accordance with section 10(a)(2) of the Federal Advisory Committee Act (5 U.S.C. App. 2), AHRQ announces meetings of the above-listed scientific peer review groups, which are subcommittees of AHRQ's Health Services Research Initial Review Group Committee. The subcommittee meetings will be closed to the public in accordance with the provisions set forth in 5 U.S.C. App. 2 section 10(d), 5 U.S.C. 552b(c)(4), and 5 U.S.C. 552b(c)(6). The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Agenda items for these meetings are subject to change as priorities dictate.

Dated: September 18, 2020.

Marquita Cullom-Stott,

Associate Director.

[FR Doc. 2020–21050 Filed 9–23–20; 8:45 am]

BILLING CODE 4160–90–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). If you wish to attend, please contact Marie Chovanec by email at *MChovanec@cdc.gov* or by telephone at 412–386–5302 at least 5 business days in advance of the meeting. She will provide you the Zoom web conference access.

DATES: The meeting will be held on November 9, 2020, from 10:00 a.m. to 2:30 p.m., EST.

ADDRESSES: This is a virtual meeting.

FOR FURTHER INFORMATION CONTACT: George W. Luxbacher, Designated Federal Officer, MSHRAC, 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 mining safety and health research projects and outcomes, including COVID–19 impact on research, funded projects, presentations, guidelines; Office of Mine Safety and Health Research (OMSHR) reshaping status; FY21 new mining projects and redesigning research; lighting research; update on MINER Act extramural research; and mining-related suicides. 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. 2020–21034 Filed 9–23–20; 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); Notice of Charter Renewal

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

ACTION: Notice of charter renewal.

SUMMARY: This gives notice under the Federal Advisory Committee Act of October 6, 1972, that the Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP), Centers for Disease Control and Prevention, Department of Health and Human Services, has been renewed for a 2-year period through September 18, 2022.

FOR FURTHER INFORMATION CONTACT: Moises C. Langub, Ph.D., Designated Federal Officer, CDC, 1600 Clifton Road NE, Mailstop TW–2, Atlanta, Georgia 30329–4027, telephone (770) 488–3585; email *MLangub@cdc.gov.*

SUPPLEMENTARY INFORMATION: 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. 2020–21035 Filed 9–23–20; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–8003, CMS–10633, CMS–10116, CMS–319, and CMS–10540]

Agency Information Collection Activities: Submission for OMB Review; 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 (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 a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate 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 on the collection(s) of information must be received by the OMB desk officer by October 26, 2020.

ADDRESSES: Written 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.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' website address at website address at *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html.*

2. Call the Reports Clearance Office at (410) 786–1326.

FOR FURTHER INFORMATION CONTACT: William Parham at (410) 786–4669.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Reinstatement with change of a previously approved collection; *Title of Information Collection:* 1915(c) Home and Community Based Services (HCBS) Waiver Application; *Use:* We will use the web-based application to review and adjudicate individual waiver actions. The web-based application will also be used by states to submit and revise their waiver requests. *Form Number:* CMS–8003 (OMB control number 0938–0449); *Frequency:* Yearly; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 47; *Total Annual Responses:* 71; *Total Annual Hours:* 6,005. (For policy questions regarding this collection contact Kathy Poisal at 410–786–5940.)

2. *Type of Information Collection Request:* Revision with change of a currently approved collection; *Title of Information Collection:* QIC Demonstration Evaluation Contractor (QDEC): Analyze Medicare Appeals to Conduct Formal Discussions and Reopenings with DME Suppliers and Part A Providers; *Use:* The Formal Telephone Discussion Demonstration and Reopenings Process is authorized under Section 402(a)(1)(F), U.S.C. 1395–1(a)(1)(F), of the Social Security Amendments of 1967. Primary and secondary data are needed to understand the effectiveness of the Demonstration in improving DME suppliers' and Part A providers' understanding of claims denial during Level 2 of the appeals process and facilitating more accurate claim submission over time. Primary data are necessary to determine, from the perspective of participating DME suppliers and Part A providers, the quality of the formal telephone discussions, satisfaction with the formal telephone discussion process, and the effect of the formal telephone discussions on submitting accurate claims. These data will inform an evaluation of the demonstration's effectiveness in achieving more accurate claims submissions, and thus reducing the number of claims CMS must process each year.

All information collected through the evaluation of the Formal Telephone Demonstration and Reopenings Process will be used by CMS through the QDEC (IMPAQ International and its partner, Palmetto GBA) to conduct analyses of satisfaction with the formal telephone discussions, and determine whether further engagement with the QIC improves understanding of the reasons for claim denials.

CMS will use the results of the evaluation to make informed policy decisions regarding the effectiveness of this demonstration and whether or not the demonstration should become a permanent part of the appeals process. Ultimately, if the information shows that DME suppliers and Part A providers were able to submit more accurate claims on the first pass, and a reduced number of claims are put through the appeals process, the Federal government could realize cost savings. *Form Number:* CMS–10633 (OMB control number: 0938–1348); *Frequency:* Yearly; *Affected Public:* Private Sector, Business or other for-profits; *Number of Respondents:* 5,288; *Total Annual Responses:* 5,288; *Total Annual Hours:* 950. (For policy questions regarding this collection contact Lynnsie G. Kelley at 410–786–1155.)

3. *Type of Information Collection Request:* Reinstatement without change of a previously approved collection; *Title of Information Collection:* Medicare Program: Conditions for Payment of Power Mobility Devices,