the comments and will use them to help disseminate and implement the plan.

Dated: March 15, 2023.

John J. Howard,

Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention.

[FR Doc. 2023–05602 Filed 3–17–23; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Center for Injury Prevention and Control

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

ACTION: Notice of meeting.

SUMMARY: In accordance with regulatory provisions, the Centers for Disease Control and Prevention (CDC) announces the following meeting for the Board of Scientific Counselors, National Center for Injury Prevention and Control (BSC, NCIPC or Board). This meeting is partially open to the public.

DATES: The meeting will be held on May 3, 2023, from 1 p.m. to 3 p.m., EDT (CLOSED); and May 4, 2023, from 9:30 a.m. to 3:30 p.m., EDT (OPEN). The public comment period will be at the end of the open session of the meeting on May 4, 2023, from 3:10 p.m. to 3:25 p.m., EDT.

ADDRESSES: Webinar, Atlanta, Georgia. All participants must register by using the following link to attend the open session: *https://cdc.zoomgov.com/meeting/register/vJIsfuqtrT4pE4H6-oCKSs9t2KHS69o3yHo.*

FOR FURTHER INFORMATION CONTACT: Christopher R. Harper, Ph.D., Senior Epidemiologist, National Center for Injury Prevention and Control, Centers for Disease Control and Prevention, 4770 Buford Highway NE, Mailstop S–1069, Atlanta, Georgia 30341. Telephone: (404) 718–8330; Email: *ncipcbsc@cdc.gov.*

SUPPLEMENTARY INFORMATION: Portions of the meeting referenced above will be closed to the public in accordance with provisions set forth in section 552b(c)(4) and (6), title 5 U.S.C., and the Determination of the Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, CDC, pursuant to Public Law 92–463 (5 U.S.C. 1009), as amended.

Purpose: The Board of Scientific Counselors, National Center for Injury Prevention and Control (BSC, NCIPC or 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 physical and mental diseases, and other impairments; (2) assist States and their political subdivisions in preventing and suppressing communicable and non-communicable diseases and other preventable conditions and in promoting health and well-being; and (3) conduct and assist in research and control activities related to injury. The BSC, NCIPC makes recommendations regarding policies, strategies, objectives, and priorities; reviews progress toward injury prevention goals; and provides evidence in injury prevention-related research and programs. The Board also provides advice on the appropriate balance of intramural and extramural research, as well as the structure, progress, and performance of intramural programs. The Board provides guidance on extramural scientific program matters, including the: (1) review of extramural research concepts for funding opportunity announcements; (2) conduct of Secondary Peer Review of extramural research grant applications, cooperative agreement applications, and contract applications received in response to funding opportunity announcements as they relate to the Center's programmatic balance and mission; (3) submission of secondary review recommendations to the Center Director of applications to be considered for funding support; (4) review of research portfolios; and (5) review of program proposals.

Matters To Be Considered: The closed session of the meeting (Day 1) will focus on the Secondary Peer Review of extramural research grant applications received in response to two (2) Notices of Funding Opportunity: RFA–CE–20–001—''Evaluating Practice-based Programs, Policies, and Practices from CDC's Rape Prevention and Education (RPE) Program: Expanding the Evidence to Prevent Sexual Violence (U01)''; and RFA–CE–23–008—''Research Grants to Develop and Validate a Prognostic Tool of Mental Health Sequelae After Traumatic Brain Injury for Adolescent Patients'' (U01).'' The open session of the meeting (Day 2) will include discussions on Improving the Quality and Reach of Extramural Research Notices of Funding Opportunity; Updated and Expanded CDC Guidance for the Identification and Response of Suicide Clusters; Developing a Cascade of Care Framework and Surveillance Indicators to Measure Linkage and Retention to Care for Substance Use Disorders; and Moving Science and Data to Violence Prevention Action. 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. 2023–05638 Filed 3–17–23; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10488]

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 May 19, 2023.

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 *https://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:_____, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, please access the CMS PRA website by copying and pasting the following web address into your web browser: *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.*

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

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

CMS–10488 Consumer Experience Survey Data Collection

Under the 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 requires Federal agencies to publish a 60-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.

Information Collection

1. *Type of Information Collection Request:* Revision of currently approved collection; *Title of Information Collection:* Consumer Experience Survey Data Collection; *Use:* Section 1311(c)(4) of the Affordable Care Act requires the Department of Health and Human Services (HHS) to develop an enrollee satisfaction survey system that assesses consumer experience with qualified health plans (QHPs) offered through an Exchange. It also requires public display of enrollee satisfaction information by the Exchange to allow individuals to easily compare enrollee satisfaction levels between comparable plans. HHS established the QHP Enrollee Experience Survey (QHP Enrollee Survey) to assess consumer experience with the QHPs offered through the Marketplaces. The survey includes topics to assess consumer experience with the health care system such as communication skills of providers and ease of access to health care services.

CMS developed the survey using the Consumer Assessment of Health Providers and Systems (CAHPS®) principles (*https://www.ahrq.gov/cahps/about-cahps/principles/index.html*) and established an application and approval process for survey vendors who want to participate in collecting QHP enrollee experience data. The QHP Enrollee Survey, which is based on the CAHPS® Health Plan Survey, will be used to (1) help consumers choose among competing health plans, (2) provide actionable information that the QHPs can use to improve performance, (3) provide information that regulatory and accreditation organizations can use to regulate and accredit plans, and (4) provide a longitudinal database for consumer research. CMS completed two rounds of developmental testing including 2014 psychometric testing and 2015 beta testing of the QHP Enrollee Survey.

The psychometric testing helped determine psychometric properties and provided an initial measure of performance for Marketplaces and QHPs to use for quality improvement. Based on psychometric test results, CMS further refined the questionnaire and sampling design to conduct the 2015 beta test of the QHP Enrollee Survey. CMS previously obtained clearance for the 2016–2023 administrations of the QHP Enrollee Survey. At this time, CMS is requesting to renew approval for the information collection related to the QHP Enrollee Experience Survey in 2024–2026. These activities are necessary to ensure that CMS fulfills legislative mandates established by section 1311(c)(4) of the Affordable Care Act to develop an ''enrollee satisfaction survey system'' and provide such information on Marketplace websites. CMS is also seeking approval to remove the flu vaccine question and revise the race and ethnicity questions to align with the 2011 HHS Data Collection Standard for the QHP Enrollee Survey 2024 administration. *Form Number:* CMS–10488 (OMB control number: 0938–1221); *Frequency:* Annually; *Affected Public Sector:* (Individuals and Households), Private Sector (Business or other for-profits and Not-for-profit institutions); *Number of Respondents:* 97,505; *Total Annual Responses:* 97,505; *Total Annual Hours:* 16,290. (For policy questions regarding this collection contact Nidhi Singh Shah at 301–492–5110).

Dated: March 14, 2023.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2023–05557 Filed 3–17–23; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers CMS–P–0015A]

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