regions/welcome-to-the-pacific-rim-region-9/land-ports-of-entry/douglas-commercial-land-port-of-entry and *https://www.gsa.gov/about-us/regions/welcome-to-the-pacific-rim-region-9/land-ports-of-entry/raul-hector-castro-land-port-of-entry.*

Serene Wetzel,

Director (Acting), Portfolio Management Division, Pacific Rim Region, Public Buildings Service.

[FR Doc. 2023–19611 Filed 9–11–23; 8:45 am]

BILLING CODE 6820–YF–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

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

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

ACTION: Notice.

SUMMARY: Notice is hereby given of a change in the meeting of the Board of Scientific Counselors, National Center for Injury Prevention and Control (BSC, NCIPC); September 20, 2023, first session from 1 p.m. to 2:30 p.m., EDT (OPEN), and second session from 2:30 p.m. to 4:30 p.m., EDT (CLOSED), in the original **Federal Register** notice.

FOR FURTHER INFORMATION CONTACT: Christopher R. Harper, Ph.D., Designated Federal Officer, Board of Scientific Counselors, 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: The virtual meeting was published in the **Federal Register** on July 31, 2023, 88 FR 49463–49464.

This meeting is being canceled in its entirety.

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. 2023–19666 Filed 9–11–23; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Notice of Closed Meeting

Pursuant to 5 U.S.C. 1009(d), notice is hereby given of the following meeting.

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

Name of Committee: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—RFA–OH–22–005, Commercial Fishing Occupational Safety Research Cooperative Agreement; and RFA–OH–22–006, Commercial Fishing Occupational Safety Training Project Grants.

Date: November 14, 2023.

Time: 1 p.m.–5 p.m., EST.

Place: Video-Assisted Meeting.

Agenda: To review and evaluate grant applications.

For Further Information Contact: Dan Hartley, Ed.D., Scientific Review Officer, Office of Extramural Programs, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, 1095 Willowdale Road, Morgantown, West Virginia 26505. Telephone: (304) 285–5812; Email: *DHartley@cdc.gov.*

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. 2023–19668 Filed 9–11–23; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10769]

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 November 13, 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 *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: ____, 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–10769 Satisfaction of Nursing Homes, Hospitals, and Outpatient Clinicians Working with the CMS Network of Quality Improvement and Innovation Contractors Program (NQIIC)

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 a currently approved information collection; *Title of Information Collection:* Satisfaction of Nursing Homes, Hospitals, and Outpatient Clinicians Working with the CMS Network of Quality Improvement and Innovation Contractors Program (NQIIC); *Use:* The purpose of this Information Collection Request (ICR) is to collect data to inform the program evaluation of the Centers for Medicare & Medicaid Services (CMS) Quality Innovation Network-Quality Improvement Organization (QIN–QIO) and Hospital Quality Improvement Contractors (HQIC) programs under the Network of Quality Improvement and Innovation Contractors (NQIIC) contract vehicle. This is a revision package. First, we updated the Nursing Home and Hospital Surveys to cover all the quality improvement focus areas targeted by NQIIC awardees, removed some but not all COVID–19 Public Health Emergency (PHE) related questions to reflect the progress of federal health program (*e.g.,* Agency for Healthcare Research and Quality Project Echo program was officially ended in August 2021), and made minor refinements based on the first round of survey fielding. Second, we added the Outpatient Clinician Survey in the same revision package since all three surveys are conducted under the same NQIIC contract.

This revision package supports evaluation of the technical assistance provided by the QINQIO Program to nursing homes and outpatient clinicians in community settings, and Hospital Quality Improvement Contractors (HQIC) Program activities to support hospitals. This ICR is part of a larger evaluation of the overall impact of the NQIIC Program. *Form Number:* CMS–10769 (OMB control number: 0938–1424); *Frequency:* Yearly; *Affected Public:* State and Private Sector (Business or other for-profits); *Number of Respondents:* 1,900; *Total Annual Responses:* 1,900; *Total Annual Hours:* 559. (For policy questions regarding this collection, contact Jeff Mokry at 214–767–4021.)

Dated: September 7, 2023.

William N. Parham, III,

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

[FR Doc. 2023–19603 Filed 9–11–23; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2023–N–0579]

Mayya Tatsene: Final Debarment Order

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is issuing an order under the Federal Food, Drug, and Cosmetic Act (FD&C Act) permanently debarring Mayya Tatsene from providing services in any capacity to a person that has an approved or pending drug product application. FDA bases this order on a finding that Ms. Tatsene was convicted of a felony under Federal law for conduct that relates to the regulation of any drug product under the FD&C Act. Ms. Tatsene was given notice of the proposed permanent debarment and was given an opportunity to request a hearing to show why she should not be debarred. As of July 8, 2023 (more than 30 days after receipt of the notice, as prescribed by regulation), Ms. Tatsene has not responded to the notice. Ms. Tatsene's failure to respond and request a hearing within the prescribed timeframe constitutes a waiver of her right to a hearing concerning this action.

DATES: This order is applicable September 12, 2023.

ADDRESSES: Any application by Ms. Tatsene for special termination of debarment under section 306(d)(4) of the FD&C Act (21 U.S.C. 335a(d)(4)) may be submitted as follows:

Electronic Submissions

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. An application submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your application will be made public, you are solely responsible for ensuring that your application does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov.*
- If you want to submit an application with confidential