and economic transportation system in the ocean commerce of the United States that is, insofar as possible, in harmony with, and responsive to, international shipping practices.'' 46 U.S.C. 40101. Pursuant to the Shipping Act, the Commission regulates ocean common carriage of the United States. When they are engaged in transportation of passengers between the U.S. and a foreign country, PVOs are common carriers under the Shipping Act. *See* 46 U.S.C. 40102(7)(A).

PVOs are also subject to the requirements of 46 U.S.C. chap. 441 and regulations promulgated thereunder in 46 CFR part 540. The purpose of that statute is, among other things, ''to prevent financial loss and hardship to the American traveling public, who, after payment of cruise passage money, are stranded by the abandonment or cancellation of a cruise.'' *Terry Marler and James Beasley dba Titanic Steamship Line,* 22 S.R.R. 359, 369 (ALJ 1983), aff'd, 22 S.R.R. 798 (FMC 1984).

The Commission understands that the current pandemic caused by the novel coronavirus (COVID–19) has severely impacted the cruise industry. On March 14, 2020, the Centers for Disease Control and Prevention (CDC) issued a No Sail Order and Suspension of Further Embarkation causing PVOs to cease all operations. Due to the unpredictable nature of this disease, the CDC has extended the term of the order demonstrating the uncertainty associated with this pandemic. Consequently, questions concerning future travel and passengers' ability to obtain refunds of monies remitted for transportation disrupted by COVID–19 are legion.

The cruise industry plays a unique and important role in the U.S. economy. Given the Commission's mandate to: (1) Ensure an efficient and economic transportation system for ocean commerce for both goods and passengers under the Shipping Act; and (2) ensure that PVOs maintain adequate financial responsibility to indemnify passengers for nonperformance and meet any liability which may be incurred for death or injury to passengers or other persons under 46 U.S.C. chap. 441, the Commission has a clear and compelling responsibility to actively investigate and respond to the current challenges impacting the cruise industry and the U.S. ports that rely on it.[1]

Therefore it is ordered, That, pursuant to 46 U.S.C. 40104, 41101–41109, 41301–41309, 44104–44106 and 46 CFR 502.281 *et seq.,* Commissioner Louis E. Sola engage cruise industry stakeholders, including PVOs, passengers, and marine terminal operators, in public or non-public discussions to identify commercial solutions to COVID–19-related issues that interfere with the operation of the cruise industry;

It is further ordered, That, the Commissioner form one or more teams, composed of leaders from the cruise industry and other stakeholders, to develop commercial solutions to the challenges created by the COVID–19 pandemic;

It is further ordered, That the Commissioner interact with any or all maritime related COVID–19 task forces of which this Commission is affiliated or monitors for the purpose of collecting data related to COVID–19 and its impact on the cruise industry;

It is further ordered, That, the Commissioner provide a preliminary report and periodic updates to the Commission on the results of efforts undertaken by this Order;

It is further ordered, That, the Commissioner have full authority under 46 CFR 502.281–291 to perform such duties as may be necessary in accordance with U.S. law and Commission regulations. The Commissioner will be assisted by staff members as may be assigned by the Chairman;

It is further ordered, That, this Proceeding be discontinued upon the acceptance of a final report and possible recommendations by the Commissioner, unless otherwise ordered by the Commission; and

It is finally ordered, That, notice of this Order be published in the **Federal Register**.

By the Commission.

Rachel Dickon,

Secretary.

[FR Doc. 2020–09623 Filed 5–5–20; 8:45 am]

BILLING CODE 6730–02–P

[1] The provisions of the Shipping Act govern proceedings under 46 U.S.C. chap. 441. *See* 46 U.S.C. 44106.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2020–0050]

Advisory Board on Radiation and Worker Health (ABRWH or the Advisory Board), National Institute for Occupational Safety and Health (NIOSH); Notice of Meeting and Request for Comment

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 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 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 24, 2020, 11:00 a.m. to 1:00 p.m., EDT.

Written comments must be received on or before June 18, 2020.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2020–0050 by any of the following methods:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* National Institute for Occupational Safety and Health, NIOSH Docket Office, 1090 Tusculum Avenue, MS C–34, Cincinnati, Ohio 45226–1998.

Instructions: All information received in response to this notice must include the agency name and docket number [CDC–2020–0050]. All relevant comments received will be posted without change to *http://www.regulations.gov,* including any personal information provided. *Instructions:* All submissions received must include the Agency name and Docket Number. All relevant comments received in conformance with the *https://www.regulations.gov* suitability policy will be posted without change to *https://www.regulations.gov,* including any personal information provided. For access to the docket to read background documents or comments received, go to *https://www.regulations.gov.*

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 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 13889 on March 22, 2020, and will terminate on March 22, 2022.

Purpose: This 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, 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.

Public Participation

Written Public Comment: The public is welcome to submit written comments in advance of the meeting. Comments should be submitted in writing by mail according to the instructions provided. The deadline for receipt of written public comment is June 18, 2020. All requests must contain the name, address, and organizational affiliation of the individual, as well as the topic being addressed. Written comments should not exceed one single-spaced typed page in length. Written comments received in advance of the meeting will be included in the official record of the meeting.

Please note that comments received, including attachments and other supporting materials, are part of the public record and are subject to public disclosure. Comments will be posted on *https://www.regulations.gov.* Therefore, do not include any information in your comment or supporting materials that you consider confidential or inappropriate for public disclosure. If you include your name, contact information, or other information that identifies you in the body of your comments, that information will be on public display. CDC will review all submissions and may choose to redact, or withhold, submissions containing private or proprietary information such as Social Security numbers, medical information, inappropriate language, or duplicate/near duplicate examples of a mass-mail campaign. CDC will carefully consider all comments submitted to the docket.

Matters to be Considered: The agenda will include discussions on: Work Group and Subcommittee Reports; Update on the Status of SEC Petitions; Plans for the August 2020 Advisory Board Meeting; and Advisory Board Correspondence. 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. 2020–09673 Filed 5–5–20; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB review; 30-Day Comment Request; Office of Minority Health Research Coordination (OMHRC) Research Training and Mentor Programs Applications (National Institute of Diabetes and Digestive and Kidney Diseases)

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below.

DATES: Comments regarding this information collection are best assured of having their full effect if received within 30-days of the date of this publication.

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.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact: Ms. Winnie Martinez, Project Officer, 6707 Democracy Blvd., 9th Floor, Bethesda, MD, 20892 or call non-toll-free number (301) 435–2988 or Email your request, including your address to: *Winnie.Martinez@nih.gov.*

SUPPLEMENTARY INFORMATION: This proposed information collection was previously published in the **Federal Register** on October 16, 2019, page 55318–55319 (84 FR 55318) and allowed 60 days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The National Institute of Diabetes and Digestive and Kidney Diseases (NIDDK), National Institutes of Health, may not conduct or sponsor, and the respondent is not required to respond to an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.