POLICIES AND PRACTICES FOR RETRIEVAL OF RECORDS:

Records will be retrieved primarily by an individual's name or business email address but may also be obtained by a search using any search term or filter.

POLICIES AND PRACTICES FOR RETENTION AND DISPOSAL OF RECORDS:

Records are retained and disposed of in accordance with FHFA's Comprehensive Record Schedule, Item 6.2 (N1–543–11–1, approved on 01/11/2013).

ADMINISTRATIVE, TECHNICAL, AND PHYSICAL SAFEGUARDS:

Electronic records are protected by controlled access procedures. Only FHFA staff (and FHFA contractors assisting such staff), whose official duties require access, are allowed to view, administer, and control these records. The System Owner controls access to this System and limits access in accordance with the above.

RECORD ACCESS PROCEDURES:

See ''Notification Procedures'' Below.

CONTESTING RECORD PROCEDURES:

See ''Notification Procedures'' Below.

NOTIFICATION PROCEDURES:

Individuals seeking notification of any records about themselves contained in this System should address their inquiry to the Privacy Act Officer, via email to *privacy@fhfa.gov* or by mail to the Federal Housing Finance Agency, 400 Seventh Street SW, Washington, DC 20219, or in accordance with the procedures set forth in 12 CFR part 1204. *Please note that all mail sent to FHFA via the U.S. Postal Service is routed through a national irradiation facility, a process that may delay delivery by approximately two weeks. For any time-sensitive correspondence, please plan accordingly.*

EXEMPTIONS PROMULGATED FOR THE SYSTEM:

None.

HISTORY:

None.

Clinton Jones,

General Counsel, Federal Housing Finance Agency.

[FR Doc. 2022–22426 Filed 10–14–22; 8:45 am]

BILLING CODE 8070–01–P

FEDERAL RETIREMENT THRIFT INVESTMENT BOARD

Notice of Meeting of the Employee Thrift Advisory Council

DATES: October 27, 2022 at 10 a.m.

ADDRESSES: Telephonic. Dial-in (listen only) information: Number: 1–202–599–1426, Code: 970 258 547#; or via web: *https://teams.microsoft.com/l/meetup-join/19%3ameeting_MmE3OGUwNTYtNTFjOC00YzQyLTg3ZjQtNmExODUwZDhiNzk5%40thread.v2/0?context=%7b%22Tid%22%3a%223f6323b7-e3fd-4f35-b43d-1a7afae5910d%22%2c%22Oid%22%3a%22533f2df2-5cef-4200-af1c-6d760c60d9cf%22%7d*

FOR FURTHER INFORMATION CONTACT: Kimberly Weaver, Director, Office of External Affairs, (202) 942–1640.

SUPPLEMENTARY INFORMATION:

Meeting Agenda

1. Approval of the minutes of the May 24, 2022, Joint Board/ETAC meeting
2. Election of ETAC Chair
3. FY22 TSP Review
4. September 2022 Investment Program Review—G, F, C, S, I, and L Funds
5. Converge Update
6. FY2023 FRTIB Budget
7. Participant Satisfaction Report
8. Legislative Update
9. New Business

Authority: 5 U.S.C. 552b(e)(1).

Dated: October 12, 2022.

Dharmesh Vashee,

General Counsel, Federal Retirement Thrift Investment Board.

[FR Doc. 2022–22512 Filed 10–14–22; 8:45 am]

BILLING CODE 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 of the Mine Safety and Health Research Advisory Committee (MSHRAC). This is a virtual meeting only. It is open to the public, limited only by the number of web conference lines (500 web conference lines are available). If you wish to attend virtually, please register according to the instructions in the addresses section below. Time will be available for public comment.

DATES: The meeting will be held on December 8, 2022, from 10:00 a.m. to 3:30 p.m., EST.

ADDRESSES: To attend the virtual meeting, please register at the website of the National Institute for Occupational Safety and Health (NIOSH) at *www.cdc.gov/niosh/mining/features/2022-12mshrac.html* or by telephone at (412) 386–4541 at least 5 business days in advance of the meeting. Registrants will receive Zoom web conference access information sent to their provided email address upon successful registration.

FOR FURTHER INFORMATION CONTACT: George W. Luxbacher, P.E., Ph.D., Designated Federal Officer, MSHRAC, NIOSH, CDC, 1600 Clifton Road NE, Mailstop V24–4, Atlanta, Georgia 30329–4027; 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 organizational structure, capabilities, projects, and outcomes. The meeting will also include an update from the NIOSH Associate Director for Mining. Agenda items are subject to change as priorities dictate.

Public Participation

Written Public Comment: The public may submit written comments or questions in advance of the meeting, to the contact person above. Written comments received in advance of the meeting will be included in the official record of the meeting, and questions will be answered during the oral public comment period open to public participation.

Oral Public Comment: This meeting will include time for members of the public to make an oral comment. The public comment session will be held on December 8, 2022, at 3:00 p.m., EST, or at the conclusion of planned presentations, and conclude following the final call for public comment. Members of the public will be given 5 to 10 minutes each for comments.

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. 2022–22454 Filed 10–14–22; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–23–1154]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled ''Generic Clearance for CDC/ATSDR Formative Research and Tool Development'' to the Office of Management and Budget (OMB) for review and approval. CDC previously published a ''Proposed Data Collection Submitted for Public Comment and Recommendations'' notice on July 22, 2022 to obtain comments from the public and affected agencies. CDC did not receive comment related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

(a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(c) Enhance the quality, utility, and clarity of the information to be collected;

(d) Minimize the burden of the collection of information on those who are to respond, including, through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses; and

(e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570. 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. Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street NW, Washington, DC 20503 or by fax to (202) 395–5806. Provide written comments within 30 days of notice publication.

Proposed Project

Generic Clearance for CDC/ATSDR Formative Research and Tool Development (OMB Control No. 0920–1154, Exp. 1/31/2023)—Extension—Office of Scientific Integrity (OSI), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention (CDC) requests approval for an Extension of a Generic Clearance for CDC/ATSDR Formative Research and Tool Development. This information collection request is designed to allow CDC to conduct formative research information collection activities used to inform many aspects of surveillance, communications, health promotion, and research project development at CDC. Formative research is the basis for developing effective strategies including communication channels, for influencing behavior change. It helps researchers identify and understand the characteristics—interests, behaviors and needs—of target populations that influence their decisions and actions.

Formative research is integral in developing programs as well as improving existing and ongoing programs. Formative research looks at the community in which a public health intervention is being or will be implemented and helps the project staff understand the interests, attributes and needs of different populations and persons in that community. Formative research occurs before a program is designed and implemented, or while a program is being conducted.

At CDC, formative research is necessary for developing new programs or adapting programs that deal with the complexity of behaviors, social context, cultural identities, and health care that underlie the epidemiology of diseases and conditions in the U.S. CDC conducts formative research to develop public-sensitive communication messages and user-friendly tools prior to developing or recommending interventions, or care. Sometimes these studies are entirely behavioral but most often they are cycles of interviews and focus groups designed to inform the development of a product.

Products from these formative research studies will be used for prevention of disease. Findings from these studies may also be presented as evidence to disease-specific National Advisory Committees, to support revisions to recommended prevention and intervention methods, as well as new recommendations.

Much of CDC's health communication takes place within campaigns that have fairly lengthy planning periods—timeframes that accommodate the standard federal process for approving data collections. Short term qualitative interviewing and cognitive research techniques have previously proven invaluable in the development of scientifically valid and population-appropriate methods, interventions, and instruments.

This request includes studies investigating the utility and acceptability of proposed sampling and recruitment methods, intervention contents and delivery, questionnaire domains, individual questions, and interactions with project staff or electronic data collection equipment. These activities will also provide information about how respondents answer questions and ways in which question response bias and error can be reduced.

This request also includes collection of information from public health programs to assess needs related to initiation of a new program activity or expansion or changes in scope or implementation of existing program activities to adapt them to current needs. The information collected will be used to advise programs and provide capacity-building assistance tailored to identify needs.

Overall, these development activities are intended to provide information that will increase the success of the surveillance or research projects through increasing response rates and decreasing response error, thereby decreasing future data collection burden to the public. The studies that will be covered under this request will include one or more of the following investigational modalities: (1) structured and qualitative interviewing for surveillance, research, interventions and