of Purpose for an Extension of Credit by a Creditor (FR T–4), and Statement of Purpose for an Extension of Credit Secured by Margin Stock (FR U–1). These reports relate to extensions of credit secured by margin stock. The Board collects the information gathered by the Margin Credit Reports so that it may meet certain obligations under the Securities Exchange Act of 1934.

Certain lenders that are not brokers, dealers, or banks making loans secured by margin stock must register and deregister with the Federal Reserve using the FR G–1 and FR G–2, respectively, and must file an annual report (FR G–4) while registered. The FR G–1, FR G–2, and FR G–4 reporting requirements collect data used to identify lenders subject to the Board's Regulation U to verify their compliance with the regulation and to monitor margin credit.

The FR T–4, FR U–1, and FR G–3 are forms that implement recordkeeping requirements for brokers and dealers, banks, and other lenders, respectively. The FR T–4 documents the purpose of credit being extended when that credit is not to purchase, carry, or trade in securities and the credit is in excess of that otherwise permitted under Regulation T. The FR G–3 and FR U–1 document the purpose of loans secured by margin stock.

Frequency: The FR G–1, FR G–2, FR G–3, FR T–4, and FR U–1 are event-generated; the FR G–4 is completed annually.

Respondents: The FR G–1, FR G–2, FR G–3, and FR G–4 panels comprise lenders, other than banks, brokers, or dealers, that extend margin credit, including federal and state credit unions; insurance companies; commercial and consumer credit organizations; production credit associations; small businesses; insurance premium funding plans; plan-lenders (a company or its affiliate that extends credit to employees to purchase company stock under an eligible employee stock option or stock purchase plan); and lenders to Employee Stock Ownership Plans (ESOPs), thrift plans, and broker-dealer affiliates. The FR T–4 panel comprises brokers and dealers and the FR U–1 panel comprises banks.

Total estimated number of respondents: FR G–1, 25; FR G–2, 12; FR G–3, 10; FR G–4, 129; FR T–4, 14; FR U–1, 14.

Estimated average hours per response: FR G–1, 1.65; FR G–2, 0.53; FR G–3, 0.25; FR G–4, 2.07; FR T–4, 0.25; FR U–1, 0.25.

Total estimated annual burden hours: 697.

Current actions: On February 16, 2024, the Board published a notice in the **Federal Register** (89 FR 12342) requesting public comment for 60 days on the extension, with revision, of the FR G–1, FR G–2, FR G–3, FR G–4, FR T–4, and FR U–1. The Board revised the FR G–1 and FR G–4 by updating the confidentiality treatment as contained in the reporting instructions to state that individual respondents may request that information submitted to the Board through the FR G–1 and FR G–4 be kept confidential and the Board will evaluate whether such treatment is appropriate on a case-by-case basis. The comment period for this notice expired on April 16, 2024. The Board did not receive any comments.

Board of Governors of the Federal Reserve System, June 25, 2024.

Benjamin W. McDonough,

Deputy Secretary and Ombuds of the Board.

[FR Doc. 2024–14341 Filed 6–27–24; 8:45 am]

BILLING CODE 6210–01–P

GENERAL SERVICES ADMINISTRATION

[Notice–MEG–2024–02; Docket No. 2024–0002; Sequence No. 29]

Notice of Establishment of a Federal Advisory Committee

AGENCY: Office of Government-wide Policy, General Services Administration (GSA).

ACTION: Notice.

SUMMARY: The General Services Administration (GSA) announces the establishment of the Open Government Federal Advisory Committee (hereinafter ''the Committee'' or ''the OG FAC'') in accordance with the Federal Advisory Committee Act, as amended.

DATES: June 28, 2024.

FOR FURTHER INFORMATION CONTACT: Arthur Brunson, Designated Federal Officer, Office of Government-wide Policy, 202–501–1126, or email: *arthur.brunson@gsa.gov;* or email: *ogfac@gsa.gov*.

SUPPLEMENTARY INFORMATION: The Administrator of the U.S. General Services Administration (GSA) established the Open Government Federal Advisory Committee (OG FAC) as a discretionary advisory committee under agency authority in accordance with the provisions of the Federal Advisory Committee Act (FACA), as amended, 5 U.S.C. 10. GSA has determined that the establishment of OG FAC is necessary and in the public interest.

GSA's Open Government Secretariat supports ensuring a more transparent, responsive and inclusive Federal Government. This is done by providing channels for members of the public to regularly engage with their government. The OG FAC will advise GSA in its endeavor to increase the public's access to data, to better advance equity, engage the public in the regulatory process, make government records more accessible, and improve the delivery of government services and benefits through expert advice.

The OG FAC will serve as an advisory body to GSA on GSA Open Government initiatives including GSA's creation, implementation and monitoring of U.S. Open Government National Action Plans (NAPs) and commitment themes. The initial focus for the OG FAC will be to provide advice to GSA on the development of NAP 6, Open Government Policy, and Public Engagement. The OG FAC will advise GSA's Administrator on emerging open government issues, challenges and opportunities to support GSA's Open Government Secretariat.

The OG FAC advisory committee is essential to conduct agency business for GSA and bring together civil society, Federal agencies, academia, industry, and other interested stakeholders. GSA needs a wide diversity of views on Open Government initiatives.

Mehul Parekh,

Acting Associate Administrator, Office of Government-wide Policy, General Services Administration.

[FR Doc. 2024–14259 Filed 6–27–24; 8:45 am]

BILLING CODE 6820–UA–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–24–24GU; Docket No. CDC–2024–0053]

Proposed Data Collection Submitted for Public Comment and Recommendations

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

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the general public and other federal agencies the opportunity to comment on

a proposed information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled Assessing Adoption and Implementation of the National Institute of Occupational Safety and Health's (NIOSH) Outputs. NIOSH proposes using surveys, interviews, and focus groups to improve awareness, understanding, and assess the impact of adoption and implementation practices by users of NIOSH research efforts and products.

DATES: CDC must receive written comments on or before August 27, 2024.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2024–0053 by either of the following methods:

- *Federal eRulemaking Portal: www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21–8, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to *www.regulations.gov.*

Please note: Submit all comments through the Federal eRulemaking portal (*www.regulations.gov*) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21–8, Atlanta, Georgia 30329; Telephone: 404–639–7570; Email: *omb@cdc.gov.*

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. In addition, the PRA also requires federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. 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;
2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
3. Enhance the quality, utility, and clarity of the information to be collected;
4. 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 submissions of responses; and
5. Assess information collection costs.

Proposed Project

Assessing Adoption and Implementation of the National Institute of Occupational Safety and Health's (NIOSH) Outputs—New—National Institute of Occupational Safety and Health's (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention (CDC), National Institute for Occupational Safety and Health (NIOSH), is requesting approval of a new Generic information collection for a period of three years for the project titled, Assessing Adoption and Implementation of the National Institute of Occupational Safety and Health's (NIOSH) Outputs.

With the continuation of the Government Performance and Results Act, and the more recent passage of the Foundations of Evidence-Based Policy Making Act, there is an increased need for federal agencies to measure and demonstrate their impact. However, measuring impact is challenging, especially for organizations that have a science-driven mission because of the time it takes to move from basic to applied research. Demonstrating attribution (cause and effect relationships) is particularly challenging for research organizations. NIOSH research is often designed to collect implementation and adoption data through document reviews of NIOSH records, including grantee final reports, and through interviews with NIOSH researchers (federal employees). While commonly recognized metrics, these data sources are not comprehensive, representative, or informative of the adoption and implementation of NIOSH products and efforts. Further, the design and execution of research projects has hindered research and program leaders prioritizing information collections to understand and assess the adoption and implementation of research efforts and products.

The proposed Generic information collection package would allow researchers to expeditiously pursue efforts to provide NIOSH with critical information to inform mission-driven needs. Additionally, the proposed efforts go beyond simply measuring customer satisfaction and seek to advance NIOSH's burden, need, and impact framework for future research while also endeavoring to execute the Office of Management and Budget's guidance regarding the Foundations of Evidence-Based Policymaking Act. Respondents are expected to consist of users and potential users of NIOSH products including subject matter expects, former NIOSH funding recipients, and intermediary and end users. CDC requests OMB approval for an estimated 6,069 annual burden hours. There is no cost to respondents other than their time to participate.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Type of data collection instrument	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Subject matter experts	Survey instrument (pre and post)	5,000	1	20/60	1,667
	Informed consent form	250	1	5/60	21
	Interview or focus group guide	250	1	1	250
Former NIOSH funding recipients	Survey instrument (pre and post)	200	1	20/60	67
	Informed consent form	25	1	5/60	2

ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Type of respondent	Type of data collection instrument	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Intermediary or end users (*e.g.,* employers, workers, manufactures, labor/professional associations, policymakers).	Interview or focus group guide	25	1	1	25
	Survey instrument (pre and post)	10,000	1	20/60	3,333
	Informed consent form	650	1	5/60	54
	Interview or focus group guide	650	1	1	650
Total					6,069

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Public Health Ethics and Regulations, Office of Science, Centers for Disease Control and Prevention.

[FR Doc. 2024–14309 Filed 6–27–24; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Committee on Breast Cancer in Young Women; 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: The Centers for Disease Control and Prevention (CDC), within the Department of Health and Human Services (HHS), announces the renewal of the charter of the Advisory Committee on Breast Cancer in Young Women (ACBCYW).

FOR FURTHER INFORMATION CONTACT: Kimberly E. Smith, M.B.A., M.H.A., Designated Federal Officer, Advisory Committee on Breast Cancer in Young Women, Centers for Disease Control and Prevention, Department of Health and Human Services, 4770 Buford Highway NE, Mailstop S107–4, Atlanta, Georgia 30341–3717. Telephone: (404) 498–0073; Email: *KESmith@cdc.gov.*

SUPPLEMENTARY INFORMATION: CDC is providing notice under 5 U.S.C. 1001–1014 of the renewal of the charter of the Advisory Committee on Breast Cancer in Young Women, Centers for Disease Control and Prevention, Department of Health and Human Services. This charter has been renewed for a two-year period through June 17, 2026.

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. 2024–14289 Filed 6–27–24; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–179, CMS–10536, CMS–R–153 and CMS–10326]

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 (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 August 27, 2024.

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