Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–23–1279]

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 ''WISEWOMAN National Program Evaluation'' 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 August 22, 2022 to obtain comments from the public and affected agencies. CDC received two comments 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

WISEWOMAN National Program Evaluation (OMB Control No. 0920–1279, Exp. 12/31/2022)—Extension—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The CDC has supported the WISEWOMAN (Well-Integrated Screening and Evaluation for Women Across the Nation) program since 1995. The WISEWOMAN program is designed to serve low-income women ages 40–64 who have elevated risk factors for cardiovascular disease (CVD) and have no health insurance, or are underinsured for medical and preventive care services. Through the WISEWOMAN program, women have access to screening services for selected CVD risk factors such as elevated blood cholesterol, hypertension, and abnormal blood glucose levels; referrals to heathy behavior support programs; and referrals to medical care. WISEWOMAN participants must be co-enrolled in the CDC-sponsored National Breast and Cervical Cancer Early Detection Program (NBCCEDP).

The WISEWOMAN program is administered through cooperative agreements with state, territorial, or tribal health departments. Each WISEWOMAN recipient submits to CDC an annual progress report that describes program objectives and activities, and semi-annual data reports (known as minimum data elements, or MDEs) on the screening, assessment, and healthy behavior support services offered to women who participate in the program. Participant-level MDEs are de-identified prior to transmission to CDC.

In 2018, CDC released the fifth funding opportunity announcement (FOA) for the WISEWOMAN program (DP18–1816), which resulted in five-year cooperative agreements with 24 state, territorial, and tribal health departments, including six new and 18 continuing awardees from the previous NOFO. Key program elements were retained (*e.g.,* provision of screening services, promotion of healthy lifestyle behaviors, and linkage to healthy behavior support services and community based resources), but a number of changes were incorporated into the program at that time. The current FOA reflects increased emphasis on three strategies to reduce CVD risk and support hypertension control and management, including: (1) tracking and monitoring clinical measures; (2) implementing team-based care; and (3) linking community resources and clinical services to support care coordination, self-management, and lifestyle change.

CDC seeks to conduct a multi-component evaluation to assess the effectiveness of the program on individual, organizational, and community-level outcomes. The in-depth assessment is designed to complement the routine progress and MDE information already being collected from WISEWOMAN program recipients. The data collection focuses on obtaining qualitative and quantitative information at the organizational and community levels about process and procedures implemented, and barriers, facilitators, and other contextual factors that affect program implementation and participant outcomes. Data collection activities include a Program Survey with all WISEWOMAN awardee programs, administered in the second and fourth program years, and a one-time site visit to each recipient spread across the three-year data collection effort. During site visits, semi-structured interviews will be conducted with WISEWOMAN staff members and staff at partner organizations, such as clinical providers and community-based resource providers, who are positioned to provide a variety of perspectives on program implementation.

CDC requests OMB approval for a one-year extension of this data collection, and requests approval for an estimated 84 annual burden hours. Participation is voluntary and there are no costs to respondents other than their time.

Estimated Annualized Burden Hours

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
WISEWOMAN Recipient Administrators	Program survey	16	1	1
	Site Visit Discussion Guide	8	1	90/60
	Innovation Site Visit Discussion Guide	2	1	45/60
Recipient partners	Site Visit Discussion Guide	16	1	1
	Innovation Site Visit Discussion Guide	2	1	45/60
Healthy behavior support staff	Site Visit Discussion Guide	16	1	1
	Innovation Site Visit Discussion Guide	2	1	45/60
Clinical providers	Site Visit Discussion Guide	16	1	1
	Innovation Site Visit Discussion Guide	2	1	45/60

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Determination Concerning a Petition To Add a Class of Employees to the Special Exposure Cohort

AGENCY: National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention, Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Department of Health and Human Services (HHS) gives notice of a determination concerning a petition to add a class of employees from the Reduction Pilot Plant in Huntington, West Virginia to the Special Exposure Cohort (SEC) under the Energy Employees Occupational Illness Compensation Program Act of 2000 (EEOICPA).

FOR FURTHER INFORMATION CONTACT: Grady Calhoun, Director, Division of Compensation Analysis and Support, National Institute for Occupational Safety and Health, 1090 Tusculum Avenue, MS C–45, Cincinnati, OH 45226–1938, Telephone: 877–222–7570. Information requests can also be submitted by email to *DCAS@CDC.GOV*.

SUPPLEMENTARY INFORMATION: On September 27, 2022, the Secretary of HHS determined that the following class of employees does not meet the statutory criteria for addition to the SEC as authorized under EEOICPA:

All International Nickel Company (INCO) security personnel who worked at any location within the Reduction Pilot Plant (RPP) during the period from June 7, 1976, through November 26, 1978.

Authority: [42 U.S.C. 7384q].

John J. Howard,

Director, National Institute for Occupational Safety and Health.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-23–23BJ; Docket No. CDC–2022–0139]

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 U.S. National Authority for Containment of Poliovirus (U.S. NAC) Data Collection Tools. Data collection will capture information relating to a poliovirus containment breach or incident at a U.S. facility and will assist the U.S. NAC in the initial stages of the investigation into the breach, and ensure that facilities have programs in place that align with global poliovirus eradication initiative.

DATES: CDC must receive written comments on or before February 10, 2023.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2022–0139 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: