STD community-based organizations; and the ethics or faith-based community. At least four members shall be persons with HIV.

Committee members are Special Government Employees (SGEs), requiring the filing of financial disclosure reports at the beginning of and annually during their terms. Individuals who are selected for appointment will be required to provide detailed information regarding their financial interests and, for example, any work they do for the federal government through research grants or contracts. Disclosure of this information is required in order for CDC ethics officials to determine whether there is a conflict between the SGE's public duties as members of CHACHSPT and their private interests, including an appearance of a loss of impartiality as defined by federal laws and regulations, and to identify any required remedial action needed to address the potential conflict. CDC and HRSA review potential candidates for CHACHSPT membership when a vacancy arises and provide a slate of nominees for consideration to the Secretary of HHS for final selection. CDC and HRSA each publishes a **Federal Register** notice and will be using a joint process to nominate nominees on a rolling basis; thus, applications received by CDC will be shared with HRSA for consideration. Therefore, potential candidates need only apply in response to one of the **Federal Register** notices. HHS notifies selected candidates of their appointment near the start of the term in December, or as soon as the HHS selection process is completed. Note that the need for different expertise varies from year to year and a candidate who is not selected in one year may be reconsidered in a subsequent year.

SGE nominees must be U.S. citizens and cannot be full-time employees of the U.S. Government. Candidates should submit the following items:

- A letter of interest or personal statement from the nominee stating how their expertise would inform the work of CHACHSPT
- A biographical sketch of the nominee (500 words or fewer)
- Current curriculum vitae or resume, including complete contact information (telephone numbers, mailing address, email address)
- At least one letter of recommendation from person(s) not employed by the U.S. Department of Health and Human Services. Candidates may submit letter(s) from current HHS employees if they wish, but at least one letter must be submitted by a person not employed by an HHS agency (*e.g.,* CDC, National Institutes of Health, Food and Drug Administration).

Nominations may be submitted directly by the individual seeking nomination or by the person/organization recommending the candidate. CDC and HRSA will collect and retain nominations received for up to two years to create a pool of potential CHACHSPT nominees. When a vacancy occurs, CDC and HRSA will review nominations and may contact nominees at that time.

Appointments shall be made without discrimination on the basis of age, race, ethnicity, gender, sexual orientation, gender identity, HIV status, disability, and cultural, religious, or socioeconomic status.

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. 2022–11815 Filed 6–1–22; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60-Day–22–0612; Docket No. CDC–2022–0074]

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 continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled Well-Integrated Screening and Evaluation for Women Across the Nation Reporting System (WISEWOMAN). The WISEWOMAN program is designed to prevent, detect, and control, hypertension and other cardiovascular disease (CVD) risk factors through services such as health coaching, and evidence informed lifestyle programs, which are tailored for individual and group behavior change.

DATES: CDC must receive written comments on or before August 1, 2022.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2022–0074 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: 404–639–7118; 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

Well-Integrated Screening and Evaluation for Women Across the Nation (WISEWOMAN) (OMB Control No. 0920–0612, Exp. 8/31/2022)—Extension—National Center for Chronic Disease and Public Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The WISEWOMAN program, sponsored by the CDC, provides services to low income, uninsured, or underinsured women aged 40–64. The WISEWOMAN program is designed to prevent, detect, and control hypertension and other cardiovascular disease (CVD) risk factors through healthy behavior support services which are tailored for individual and group behavior change. The WISEWOMAN program provides services to women who are jointly enrolled in the National Breast and Cervical Cancer Early Detection Program (NBCCEDP), which is also administered by CDC.

The WISEWOMAN program is administered by state health departments and tribal programs. In 2018, a new five-year cooperative agreement was awarded under Funding Opportunity Announcement DP18–1816, subject to the availability of funds. CDC collects two types of information from WISEWOMAN awardees, which is submitted in an electronic data file to CDC twice per year. The Minimum Data Elements (MDE) file contains data using a unique identifier with client-level information about CVD risk factors and types of healthy behavior support services for participants served by the program. The estimated burden per response for the MDE file is 24 hours. The Annual Progress Report provides a narrative summary of each awardee's objectives and the activities undertaken to meet program goals. The estimated burden per response is 16 hours.

There are no changes to the information collected. CDC will continue to use the information collected from WISEWOMAN awardees to support program monitoring and improvement activities, evaluation, and assessment of program outcomes. The overall program evaluation helps to demonstrate program accomplishments and strengthen the evidence for strategy implementation for improved engagement of underserved populations. It can also determine whether the identified strategies and associated activities can be implemented at various levels within a state or tribal organization. Evaluation is also designed to demonstrate how WISEWOMAN can obtain cardiovascular disease health outcome data on at-risk populations, promote public education about CVD risk-factors, and improve the availability of healthy behavior support services for under-served women.

CDC requests a two-year Extension of this data collection. The total estimated annual burden hours are 2,240. Participation in this information collection is required as a condition of cooperative agreement funding. 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	Aervage burden per response (in hrs.)	Total burden (in hrs.)
WISEWOMAN Awardees	Screening and Assessment and Lifestyle Program MDEs.	35	2	24	1,680
	Annual Progress Report	35	1	16	560
Total					2,240

Jeffrey M. Zirger,

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

[FR Doc. 2022–11772 Filed 6–1–22; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–22–22FS; Docket No. CDC–2022–0071]

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 Artificial Stone Countertops: Exposures, Controls, Surveillance, & Translation. The purpose of the proposed data collection is to conduct a survey with artificial stone countertop fabrication facilities to better understand, work practices and controls related to respirable crystalline silica, barriers or facilitators to implementation of medical and exposure monitoring requirements, and