

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–22–0612]

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 “Well-Integrated Screening and Evaluation for Women Across the Nation (WISEWOMAN) Reporting System” 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 June 2, 2022 to obtain comments from the public and affected agencies. CDC received no comment(s) 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

Well-Integrated Screening and Evaluation for Women Across the Nation (WISEWOMAN) Reporting System (OMB Control No. 0920–0612, Exp. 08/31/2022)—Extension—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The WISEWOMAN program, sponsored by the CDC, provides services to low income, uninsured, or underinsured women aged 40–64. WISEWOMAN 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 sponsored by CDC.

The WISEWOMAN program is administered by state health departments and tribal programs. In 2018, new five-year cooperative agreements were awarded under

Funding Opportunity Announcement DP18–1816, subject to the availability of funds. CDC collects two types of information from WISEWOMAN awardees. The WISEWOMAN awardee submits 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 cardiovascular disease 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. In addition, each WISEWOMAN awardee submits an Annual Progress Report to CDC, which provides a narrative summary of the awardee’s objectives and the activities undertaken to meet program goals. The estimated burden per response for the Annual Progress Report is 16 hours.

There are no changes to the information collection. 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 implementing strategies that improve engagement of underserved populations. The information reported to CDC can also help to 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 cardiovascular disease risk-factors, and improve the availability of healthy behavior support services for under-served women.

OMB approval is requested for two years. Participation in this information collection is required as a condition of cooperative agreement funding. There are no costs to respondents other than their time. The total estimated annualized burden hours are 2,240.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)
WISEWOMAN Awardees	Screening and Assessment and Lifestyle Program MDEs.	35	2	24
	Annual Progress Report	35	1	16

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-22-0770]

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 “National HIV Behavioral Surveillance System (NHBS)” 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 May 13, 2022 to obtain comments from the public and affected agencies. CDC did not receive 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

National HIV Behavioral Surveillance System (NHBS) (OMB Control No. 0920-0770, Exp. 01/31/2023)—Revision—National Center for HIV, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The purpose of this data collection is to monitor behaviors of persons at high risk for infection that are related to Human Immunodeficiency Virus (HIV) transmission and prevention in the United States. The primary objectives of the NHBS are to obtain data from samples of persons at risk to: (a) describe the prevalence and trends in risk behaviors; (b) describe the prevalence of and trends in HIV testing and HIV infection; (c) describe the prevalence of and trends in use of HIV prevention services; and (d) identify met and unmet needs for HIV prevention services in order to inform health departments, community based organizations, community planning groups and other stakeholders.

By describing and monitoring the HIV risk behaviors, HIV seroprevalence and incidence, and HIV prevention experiences of persons at highest risk for HIV infection, NHBS provides an important data source for evaluating progress towards national public health initiatives, such as reducing new infections, increasing the use of

condoms, and targeting populations at high risk.

The Centers for Disease Control and Prevention requests approval for a three-year Revision of this information collection. Data are collected through in-person interviews conducted with persons systematically selected from 20 Metropolitan Statistical Areas (MSAs) throughout the United States; these 20 MSAs are chosen based on highest number of HIV infections diagnosed. Persons at risk for HIV infection to be interviewed for NHBS include men who have sex with men (MSM), persons who inject drugs (PWID), and heterosexually active persons at increased risk of HIV infection (HET). A brief screening interview will be used to determine eligibility for participation in the behavioral assessment.

The data from the behavioral assessment will provide estimates of: (1) behavior related to the risk of HIV and other sexually transmitted diseases; (2) prior testing for HIV; and (3) use of HIV prevention services.

All persons interviewed will also be offered an HIV test and will participate in a pre-test counseling session. No other federal agency systematically collects this type of information from persons at risk for HIV infection. These data have substantial impact on prevention program development and monitoring at the local, state, and national levels. CDC estimates that NHBS will involve, per year, in up to 20 MSAs, eligibility screening for 125 persons and eligibility screening plus the behavioral assessment with 500 eligible respondents, resulting in a total of 30,000 eligible survey respondents and 7,500 ineligible screened persons during a three-year period. Data collection will rotate such that interviews will be conducted among one group per year: MSM in Year 1, PWID in Year 2, and HET in Year 3. The type of data collected for each group will vary slightly due to different sampling methods and risk characteristics of the group.

Participation of respondents is voluntary and there is no cost to the respondents other than their time. CDC requests OMB approval for an estimated 6,600 annual burden hours.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Persons Screened	Eligibility Screener	12,500	1	5/60
Eligible Participants	Behavioral Assessment MSM	3,333	1	24/60