

ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hr)
Child MUH Residents in LA County MUH Residents in Minnesota, Maine and Florida.	Resident Survey—Post Intervention: Core	500	1	45/60
	Resident Survey—Post Intervention: Children's Module.	250	15/60
	Protocol for Saliva Collection (Adult)	1,000	1	10/60
	Airborne Particle Monitoring Diary	200	1	90/60
	Protocol for Saliva Collection (Child)	500	1	10/60
	Resident Focus Group Telephone Screening Interview Script.	60	1	5/60
	Resident Pre-Focus Group Demographic and Attitudinal Survey.	60	1	5/60
	MUH Resident Focus Group Guide—Process Oriented.	30	1	1
	MUH Resident Focus Group Guide—Outcome Oriented.	30	1	1

Dated: October 2, 2012.

Ron A. Otten,

Director, Office of Scientific Integrity (OSI)
Office of the Associate Director for Science
(OADS), Office of the Director, Centers for
Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30-Day–13–12SF]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639-7570 or send an email to omb@cdc.gov. Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395-5806. Written comments should be received within 30 days of this notice.

Proposed Project

Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery—NEW—Centers for Disease Control and Prevention (CDC), National Institute for Occupational Safety and Health (NIOSH).

As part of a Federal Government-wide effort to streamline the process to seek feedback from the public on service

delivery, the CDC has submitted a Generic Information Collection Request (Generic ICR): “Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery” to OMB for approval under the Paperwork Reduction Act (PRA) (44 U.S.C. 3501 *et seq.*).

To request additional information, please contact Kimberly S. Lane, Reports Clearance Officer, Centers for Disease Control and Prevention, 1600 Clifton Road, MS-D74, Atlanta, GA 30333 or send an email to omb@cdc.gov.

SUPPLEMENTARY INFORMATION:

Title: Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery.

Abstract: The information collection activity will garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Administration's commitment to improving service delivery. By qualitative feedback we mean information that provides useful insights on perceptions and opinions, but are not statistical surveys that yield quantitative results that can be generalized to the population of study. This feedback will provide insights into customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative and actionable communications between the Agency and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

Feedback collected under this generic clearance will provide useful information, but it will not yield data that can be generalized to the overall population. This type of generic clearance for qualitative information will not be used for quantitative information collections that are designed to yield reliably actionable results, such as monitoring trends over time or documenting program performance. Such data uses require more rigorous designs that address: The target population to which generalizations will be made, the sampling frame, the sample design (including stratification and clustering), the precision requirements or power calculations that justify the proposed sample size, the expected response rate, methods for assessing potential non-response bias, the protocols for data collection, and any testing procedures that were or will be undertaken prior fielding the study. Depending on the degree of influence the results are likely to have, such collections may still be eligible for submission for other generic mechanisms that are designed to yield quantitative results.

The Agency received no comments in response to the 60-day notice published in the **Federal Register** on December 22, 2010 (75 FR 80542).

This is a new collection of information. Respondents will be screened and selected from Individuals and Households, Businesses, Organizations, and/or State, Local or Tribal Government. Below we provide CDC's projected annualized estimate for the next three years. There is no cost to respondents other than their time. The estimated annualized burden hours for this data collection activity are 28,750.

Type of collection	Average number of respondents per activity	Annual frequency per response	Average number of activities	Average hours per response
Online surveys, Telephone Surveys, Focus Groups, In person observation/testing	14,350	1	4	30/60

Dated: October 2, 2012.

Ron A. Otten,

Director, Office of Scientific Integrity (OSI), Office of the Associate Director for Science (OADS), Office of the Director, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-13-12MQ]

Agency Forms Undergoing Paperwork Reduction Act Review

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Proposed Project

Evaluation of the Young Sisters Initiative: A Guide to A Better You! Program—New—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

In 2010, the Centers for Disease Control and Prevention (CDC) launched

the three-year Breast Cancer in Young Women (BCYW) project to raise awareness about these issues among young breast cancer survivors (YBCS) and to provide psychosocial and reproductive health support to women who are diagnosed before age 45. A key component of the BCYW program is the design, testing, implementation and evaluation of the Young Sisters Initiative: A Guide to a Better You (YSI) program. The YSI program is a web-based intervention designed to provide African American YBCS with culturally tailored psychosocial and reproductive health information to support their needs as cancer survivors.

CDC plans to conduct a process evaluation of YSI program implementation in conjunction with Sisters Network Inc. (SNI), a partner organization, and ICF International, an evaluation contractor. Information will be collected to assess whether the YSI program can be implemented with fidelity; reach its target audience of African American YBCS; and deliver effective psychosocial and reproductive health information and support. The process evaluation will also collect information to improve understanding of facilitators and barriers to YSI program recruitment and implementation, and to assess how the program might be adapted for use with other audiences.

Primary information collection will consist of two Web-based surveys of YSI program users, conducted before and after exposure to YSI program materials. The initial five-minute demographic screener will be conducted when users encounter the YSI Web site. Respondents will be asked to provide demographic and health information necessary for identifying members of the

target YSI program audience, and to indicate their willingness to complete a brief online post-use survey one to two weeks after their initial YSI program Web site visit. The post-use survey will be conducted after YSI Web site users have time to review the site and materials. The estimated burden for the post-use survey is 20 minutes. Respondents will be asked questions about the usefulness of resources posted on the YSI Web site and satisfaction with the site. No personally identifiable information will be collected.

Two secondary sources of information will be used to supplement the process evaluation data collection, but will not impose burden on YSI Web site users. First, CDC's evaluation contractor will use information obtained through Google Analytics to assess how visitors (particularly the target audience) navigate and use the YSI Web site. In addition, the evaluation contractor will conduct a limited number of telephone interviews with SNI staff and SNI-identified recruitment partners before and after the YSI implementation to assess fidelity to the YSI program core components and identify any facilitators and/or barriers experienced during program implementation.

CDC will use the results of the process evaluation to inform future efforts to support and educate YBCS in vulnerable/minority populations. OMB approval is requested for one year. Participation in the information collection is voluntary, and there are no costs to respondents other than their time. The total estimated annualized burden hours are 142.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hr)
YSI Web Site Users	YSI Program Demographic Screener	500	1	5/60
	YSI Program Post-Use Survey	300	1	20/60