

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–15–0696; Docket No. CDC–2015–0022]

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 efforts to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. This notice invites comment on the information collection request entitled *National HIV Prevention Program Monitoring and Evaluation (NHM&E)*. CDC is requesting a 3-year approval for revision to the previously approved project to continue collecting standardized HIV prevention program evaluation data from health departments and community-based organizations (CBOs) who receive federal funds for HIV prevention activities.

DATES: Written comments must be received on or before June 22, 2015.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2015–0022 by any of the following methods:

- *Federal eRulemaking Portal:* Regulations.gov. Follow the instructions for submitting comments.

- *Mail:* Leroy A. Richardson, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS–D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. All relevant comments received will be posted without change to Regulations.gov, including any personal information provided. For access to the docket to read background documents or comments received, go to Regulations.gov.

Please note: All public comment should be submitted through the Federal eRulemaking portal (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 the Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS–D74, Atlanta, Georgia 30329; phone: 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 OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology; and (e) estimates of capital or start-up costs and costs of operation, maintenance, and purchase of services to provide information. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information; and to transmit or otherwise disclose the information.

Proposed Project

National HIV Prevention Program Monitoring and Evaluation (NHM&E) (OMB 0920–0696, Expiration 03/31/2016)—Revision—National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

CDC is requesting a 3-year approval for revision to the previously approved project. The purpose of this revision is to continue collecting standardized HIV prevention program evaluation data from health departments and community-based organizations (CBOs) who receive federal funds for HIV prevention activities. Grantees have the option of key-entering or uploading data to a CDC-provided web-based software application (EvaluationWeb®).

This revision includes changes to the data variables to adjust to the different monitoring and evaluation needs of new funding announcements without a change in burden.

The evaluation and reporting process is necessary to ensure that CDC receives standardized, accurate, thorough evaluation data from both health department and CBO grantees. For these reasons, CDC developed standardized NHM&E variables through extensive consultation with representatives from health departments, CBOs, and national partners (e.g., The National Alliance of State and Territorial AIDS Directors, Urban Coalition of HIV/AIDS Prevention Services, and National Minority AIDS Council).

CDC requires CBOs and health departments who receive federal funds for HIV prevention to report non-identifying, client-level and aggregate-level, standardized evaluation data to: (1) Accurately determine the extent to which HIV prevention efforts are carried out, what types of agencies are providing services, what resources are allocated to those services, to whom services are being provided, and how these efforts have contributed to a reduction in HIV transmission; (2) improve ease of reporting to better meet these data needs; and (3) be accountable to stakeholders by informing them of HIV prevention activities and use of funds in HIV prevention nationwide.

CDC HIV prevention program grantees will collect, enter or upload, and report agency-identifying information, budget data, intervention information, and client demographics and behavioral risk characteristics with an estimate of 200,846 burden hours. Data collection will include searching existing data sources, gathering and maintaining data,

document compilation, review of data, and data entry or upload into the web-based system.

There are no additional costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Health jurisdiction	Health Department Reporting	69	2	1,377	190,026
Community-Based Organization	Community-Based Organization Reporting.	200	2	40.5	16,200
Total	206,226

Leroy A. Richardson,
Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.
 [FR Doc. 2015-09088 Filed 4-20-15; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

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[30Day-15-0314]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (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 or send an email to *omb@cdc.gov*. Written comments and/or suggestions regarding the items contained in this notice should be directed to the Attention: 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

The National Survey of Family Growth (NSFG)-(0920-0314, Expiration 04/30/2015—Revision—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Section 306 of the Public Health Service (PHS) Act (42 U.S.C. 242k), as amended, authorizes that the Secretary of Health and Human Services (DHHS), acting through NCHS, shall collect statistics on “family formation, growth, and dissolution,” as well as “determinants of health” and “utilization of health care” in the United States. This three-year clearance request includes the data collection in 2015–2018 for the continuous NSFG.

The National Survey of Family Growth (NSFG) was conducted periodically between 1973 and 2002, and continuously since 2006, by the National Center for Health Statistics, CDC. Each year, about 15,000 households are screened, with about 5,000 participants interviewed annually. Participation in the NSFG is completely voluntary and confidential. Interviews average 60 minutes for males and 80 minutes for females. The response rate since 2011 has been about 73 percent.

The NSFG program produces descriptive statistics which measure factors associated with birth and pregnancy rates, including contraception, infertility, marriage, divorce, and sexual activity, in the U.S. population 15–49; and behaviors that affect the risk of sexually transmitted diseases (STD), including HIV, and the medical care associated with contraception, infertility, and pregnancy and childbirth.

NSFG data users include the DHHS programs that fund it, including CDC/NCHS and nine others (The Eunice Kennedy Shriver National Institute of Child Health and Human Development (NIH/NICHHD); the Office of Population Affairs (DHHS/OPA); the Children’s Bureau within the Administration for Children and Families (DHHS/ACF/CB); the ACF’s Office of Planning, Research, and Evaluation (DHHS/ACF/OPRE); the CDC’s Division of HIV/AIDS Prevention (CDC/DHAP); the CDC’s Division of STD Prevention (CDC/DSTDP); the CDC’s Division of Cancer Prevention and Control (CDC/DCPC); the CDC’s Division of Birth Defects and Developmental Disabilities (CDC/DBDDD); and the CDC’s Division of Reproductive Health (CDC/DRH). The NSFG is also used by state and local governments; private research and action organizations focused on men’s and women’s health, child well-being, and marriage and the family; academic researchers in the social and public health sciences; journalists, and many others.

No questionnaire changes are requested in the first 6 months of this clearance; limited changes including (1) the expansion of the age range from 15–44 years of age to 15–49, (2) some revision of the female and male questionnaires to incorporate new and modified items related to contraceptive use, reproductive health, preventive service screening/counseling, sexual orientation, health insurance, cigarette smoking, cancer risk, military service