

EXHIBIT 1—ESTIMATED BURDEN HOURS FOR THE 2015 LONGITUDINAL SURVEY

Form name	Number of respondents	Number of responses per respondent	Hours per response	Total burden hours
Prescreener Questionnaire	4,300	1	0.09	387
Establishment Questionnaire	2,054	1	* 0.38	781
Plan Questionnaire	2,054	1.4	0.18	518
Total	8,408	na	na	1,686

* The burden estimate printed on the establishment questionnaire is 45 minutes which includes the burden estimate for completing the establishment questionnaire, an average of 1.4 plan questionnaires, plus the prescreener. The establishment and plan questionnaires are sent to the respondent as a package and are completed by the respondent at the same time.

EXHIBIT 2—ESTIMATED COST BURDEN FOR THE 2015 LONGITUDINAL SURVEY

Form name	Number of respondents	Total burden hours	Average hourly wage rate*	Total burden hours
Prescreener Questionnaire	4,300	387	\$30.44	\$11,780
Establishment Questionnaire	2,054	781	\$30.44	\$23,774
Plan Questionnaire	2,054	518	\$30.44	\$15,768
Total	8,408	1,686	na	\$51,322

*Based upon the mean hourly wage for Compensation, Benefits, and Job Analysis Specialists occupation code 13-1141, at <http://www.bls.gov/oes/current/oes131141.htm> (U.S. Department of Labor, Bureau of Labor Statistics).

Request for Comments

In accordance with the Paperwork Reduction Act, comments on AHRQ's information collection are requested with regard to any of the following: (a) Whether the proposed collection of information is necessary for the proper performance of AHRQ health care research and information dissemination functions, including whether the information will have practical utility; (b) the accuracy of AHRQ's estimate of burden (including hours and costs) of the proposed collection(s) of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information upon the respondents, including the use of automated collection techniques or other forms of information technology.

Comments submitted in response to this notice will be summarized and included in the Agency's subsequent request for OMB approval of the proposed information collection. All comments will become a matter of public record.

Dated: February 5, 2015.

Richard Kronick, Ph.D.,

AHRQ Director.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES**Centers for Disease Control and Prevention**

[30Day-15-0920]

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

Data Collection Through Web Based Surveys for Evaluating Act Against AIDS Social Marketing Campaign Phases Targeting Consumers (Generic ICR, OMB# 0920-0920, Expires 2/28/2015)—Extension—National Center for HIV/AIDS, Viral Hepatitis, STD and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

In response to the continued HIV epidemic in our country, CDC has launched Act Against AIDS, a 5-year, multifaceted communication campaign to reduce HIV incidence in the United States. CDC plans to release the campaign in phases, with some of the phases running concurrently. Each phase of the campaign will use mass media and direct-to-consumer channels to deliver HIV prevention and testing messages. Some components of the campaign will be designed to provide

basic education and increase awareness of HIV/AIDS among the general public, and others will be targeted to specific subgroups or communities at greatest risk of infection. The current study addresses the need to assess the effectiveness of these social marketing messages aimed at increasing HIV awareness and delivering HIV prevention and testing messages among at-risk populations.

This extension of an ongoing study will evaluate the *Act Against AIDS* (AAA) social marketing campaign aimed at increasing HIV/AIDS awareness, increasing prevention behaviors, and improving HIV testing rates among consumers. A total of 36,000 respondents were originally approved for this 3-year data collection. Since the

original approval date, 4,250 respondents have participated in the surveys. The number of remaining respondents for the 3-year period is 31,750. We anticipate screening approximately 52,915 individuals annually to achieve 10,583 respondents annually. The information collected from each of the data collections were used to evaluate specific AAA campaign phases. We are requesting additional time to continue to survey other AAA target audiences and campaign phases and measuring exposure to each phase of the campaign and interventions implemented under AAA.

Depending on the target audience for the campaign phase, the study screener will vary. The study screener may address one or more of the following

items: race/ethnicity, sexual behavior, and sexual orientation. Each survey will have a core set of items asked in all rounds, as well as a module of questions relating to specific AAA activities and communication initiatives.

Survey respondents will be selected from a combination of sources, including a national opt-in email list sample and respondent lists generated by partnership organizations (e.g., the National Urban League, the National Medical Association). Participants will self-administer the survey at home on personal computers. There is no cost to the respondents other than their time. The total number of estimated annual burden hours is 7,056.

ESTIMATED ANNUALIZED BURDEN HOURS

Respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Individuals (male and female) aged 18 years and older/Study Screener.	Study Screener	52,915	1	2/60
Individuals (male and female) aged 18 years and older.	Survey	10,583	1	30/60

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.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-15-0010]

Proposed Data Collections Submitted for Public Comment and Recommendations

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 to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. To request more information on the below proposed project or to obtain a copy of the information collection plan and instruments, call 404-639-7570 or send

comments to Leroy A. Richardson, 1600 Clifton Road, MS-D74, Atlanta, GA 30333 or send an email to omb@cdc.gov.

Comments submitted in response to this notice will be summarized and/or included in the request for Office of Management and Budget (OMB) approval. 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. Written comments should be received within 60 days of this notice.

Proposed Project

Birth Defects Study To Evaluate Pregnancy exposures (BD-STEPS) (formerly titled The National Birth Defects Prevention Study (NBDPS)), (OMB 0920-0010, Expiration 01/31/2017)—Revision—National Center on Birth Defects and Developmental Disabilities (NCBDDD), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

CDC has been monitoring the occurrence of serious birth defects and genetic diseases in Atlanta since 1967 through the Metropolitan Atlanta Congenital Defects Program (MACDP). The MACDP is a population-based surveillance system for birth defects currently covering three counties in Metropolitan Atlanta.

Since 1997, CDC has funded case-control studies of major birth defects that utilize existing birth defect surveillance registries (including