

ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Customer comment cards, interactive voice surveys	GenIC_Request Template	61,000	1	15/60

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 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**

[30Day-17-16AWJ]

**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

Behavioral Risk Factor Surveillance System (BRFSS) Asthma Call-back Survey (ACBS)—Existing Collection in Use without an OMB Control Number—National Center for Environmental Health (NCEH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention (CDC) is requesting a three-year Paperwork Reduction Act (PRA) clearance to conduct information collection under “The Behavioral Risk Factor Surveillance System (BRFSS) Asthma Call-back Survey (ACBS)” for three years beginning with the 2017 data collection cycle. The ACBS is an existing collection in use without an OMB Control Number. BRFSS (OMB Control No. 0920-1061, expiration date 3/31/2018) is a nationwide system of customized, cross-sectional telephone health surveys sponsored by CDC’s National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP) Division of Population Health. The BRFSS information collection is conducted in a continuous, three-part telephone interview process: Screening, participation in a common BRFSS core survey, and participation in optional question modules that states use to customize survey content.

The ACBS is not an optional state module, but rather, is a follow-up survey to the regular BRFSS efforts. It is funded by the National Asthma Control Program (NACP) in the Air Pollution and Respiratory Health Branch (APRHB) of the National Center for Environmental Health (NCEH). The ACBS is administered by NCCDPHP on behalf of NCEH using its existing BRFSS sampling frame. BRFSS coordinators in the health departments in U.S. states, territories, and the District of Columbia

(collectively referred to as states) are responsible for survey administration. Currently CDC provides its 40 participating states with technical and methodological assistance.

The purpose of ACBS is to gather state-level asthma data and to make them available to track the burden of the disease, to monitor adherence to asthma guidelines, and to direct and evaluate interventions undertaken by asthma control programs located in state health departments. Beyond asthma prevalence estimates, for most states, the ACBS provides the only sources of adult and child asthma data on the state and local level.

As a follow-up, the ACBS is conducted within two weeks after the BRFSS survey. Data collection for ACBS involves (1) screening, (2) obtaining permission, (3) consenting and telephone interviewing on a subset of the BRFSS respondents from participating states. The ACBS eligible respondents are BRFSS adults, 18 years and older, who report ever being diagnosed with asthma. In addition, some states include children, below 18 years of age, who are randomly selected subjects in the BRFSS household. Parents or guardians serve as ACBS proxy respondents for their children ever diagnosed with asthma. If both the BRFSS adult respondent and the selected child in the household have asthma, then only one or the other is eligible for the ACBS.

The ACBS adds considerable state-level depth to the existing body of asthma data. It addresses critical questions surrounding the health and experiences of persons with asthma. Health data include symptoms, environmental factors, and medication use among persons with asthma. Data on their experiences include activity limitation, health system use, and self-management education. These asthma data are needed to direct and evaluate interventions undertaken by asthma control programs located in state health departments. Federal agencies and other entities also rely on this critical information for planning and evaluating efforts and to reduce the burden from this disease.

The CDC makes annual ACBS datasets available for public use and provides

guidance on statistically appropriate uses of the data. Participation in the ACBS is voluntary and there are no costs to respondents other than their time. The burden table reflects the landline and cell phone data collection

methods used in 2013 and later years. Additionally, the burden table accounts for reporting burden incurred by the states for the monthly or quarterly data submission to CDC. The burden hour estimates represent the 2013 data

collection which is the most recent data released.

The total estimated annualized burden hours for all respondents are 6,029 hours.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)
BRFSS Adults	ACBS Landline Screener—Adult	21,424	1	1/60
	ACBS Cell Phone Screener—Adult	8,976	1	1/60
BRFSS Parents or Guardians of Children	ACBS Landline Screener—Child	4,245	1	1/60
	ACBS Cell Phone Screener—Child	2,238	1	1/60
ACBS Adults	ACBS Adult Consent and Survey—2013	19,954	1	10/60
ACBS Parents or Guardians of Children	ACBS Child Consent and Survey—2013	3,887	1	10/60
State BRFSS Coordinators	ACBS Data Submission Layout	40	12	3

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: Accomplishments of the Domestic Violence Hotline, Online Connections and Text (ADVHOCaT) Study.

OMB No.: 0970-0468.

Description: The National Domestic Violence Hotline (The Hotline) and

loveisrespect (LIR), which are supported by the Division of Family Violence Prevention and Services within the Family and Youth Services Bureau (FYSB) of the Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), serve as partners in the intervention, prevention, and resource assistance efforts of the network of domestic violence and dating violence service providers.

In order to describe the activities and accomplishments of The Hotline and LIR and develop potential new or revised performance measures, the ACF/HHS Office of Planning, Research and Evaluation (OPRE) and FYSB are proposing a data collection activity as part of the Accomplishments of the Domestic Violence Hotline, Online Connections and Text (ADVHOCaT) Study.

As part of ongoing program activities and monitoring for The Hotline and LIR,

ACF proposes to collect information via voluntary phone, chat, and web-based surveys of individuals who contact The Hotline and LIR. Participants will complete a baseline survey at the end of their contact with The Hotline and LIR, and a follow-up survey approximately two weeks later. The survey will include questions about reasons for contacting The Hotline/LIR, whether needs were met, satisfaction with services received, and helpfulness of information provided. This data collection builds on a previous data collection that was focused on understanding the preferred mode of contact by those who contact The Hotline and LIR. This new information will inform future efforts to monitor and improve the performance of domestic violence hotlines and provide hotline services.

Respondents: Individuals aged 18 and older who contact The Hotline and LIR via phone or chat.

ANNUAL BURDEN ESTIMATES—2 YEAR REQUEST

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Annual burden hours
The Hotline/LIR Baseline Survey	2200	1100	1	0.056	62
The Hotline/LIR Follow Up Survey	2200	1100	1	0.1	110

Estimated Total Annual Burden Hours: 172.

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and

comments may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW., Washington, DC 20201, Attn: OPRE Reports Clearance Officer. Email address: OPREinfocollection@acf.hhs.gov. All requests should be identified by the title of the information collection.

The Department specifically requests comments 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) the quality, utility, and clarity of the information to be collected; and (d)