

the second leading cause of death from cancer in the United States. CRC screening has been shown to reduce incidence of and death from the disease. Screening for CRC can detect disease early when treatment is more effective and prevent cancer by finding and removing precancerous polyps. Of individuals diagnosed with early stage CRC, more than 90% live five or more years. Despite strong evidence supporting screening, only 65% of adults currently report being up-to-date with CRC screening as recommended by the U.S. Preventive Services Task Force, with more than 22 million age-eligible adults estimated to be untested. To reduce CRC morbidity, mortality, and associated costs, use of CRC screening tests must be increased among age-eligible adults with the lowest CRC screening rates.

CDC's Colorectal Cancer Control Program (CRCCP) currently provides funding to 31 grantees under "Organized Approaches to Increase Colorectal Cancer Screening" (CDC–RFA–DP15–1502). CRCCP grantees include state governments or bona-fide agents, universities, and tribal organizations. The purpose of the new cooperative agreement program is to

increase CRC screening rates among an applicant defined target population of persons 50–75 years of age within a partner health system serving a defined geographical area or disparate population.

The CRCCP was significantly redesigned in 2015 and has two components. Under Component 1, all 31 CRCCP grantees receive funding to support partnerships with health systems to implement up to four priority evidence-based interventions (EBIs) described in the Guide to Community Preventive Services, as well as other supporting strategies. Grantees must implement at least two EBIs in each partnering health system. Under Component 2, 6 of the 31 CRCCP grantees will provide direct screening and follow-up clinical services for a limited number of individuals aged 50–64 in the program's priority population who are asymptomatic, at average risk for CRC, have inadequate or no health insurance for CRC screening, and are low income

Based on the redesigned CRCCP, the information collection plan has also been redesigned to address the two program components. The new cooperative agreement program (CDC–

RFA–DP15–1502) requires that CDC monitor and evaluate the CRCCP and individual grantee performance using both process and outcome evaluation. Two forms of data collection are proposed. First, the CRCCP grantee survey was redesigned to align with new CRCCP goals. The grantee survey will be submitted to CDC annually. Second, CDC proposes to collect clinic-level data to assess changes in CDC's primary outcome of interest, *i.e.*, CRC screening rates within partner health systems. Each grantee will complete a clinic-level data template once per month. All information will be reported to CDC electronically.

The information collection will enable CDC to gauge progress in meeting CRCCP program goals and to monitor implementation activities, evaluate outcomes, and identify grantee technical assistance needs. In addition, findings will inform program improvement and help identify successful activities that need to be maintained, replicated, or expanded.

OMB approval is requested for three years. Participation is required for CRCCP awardees. There are no 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 hr)	Total burden (in hr)
CRCCP Grantees	CRCCP Annual Grantee Survey.	31	1	45/60	23
	CRCCP Clinic-level Data Collection Template.	31	12	30/60	186
Total	209

Leroy A. Richardson,
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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

[60 Day–16–1061; Docket No. CDC–2016–0008]

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 Behavioral Risk Factor Surveillance System (BRFSS), a state-level survey of health risk behaviors and chronic health conditions. Survey questions are updated each year. The information collection is being revised to incorporate an annual field test of proposed changes prior to their implementation on a broad scale.

DATES: Written comments must be received on or before March 21, 2016.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2016–0008 by any of the following methods:

Federal eRulemaking Portal: Regulation.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

Behavioral Risk Factor Surveillance System (BRFSS) (OMB No. 0920-1061, exp. 3/31/2018)—Revision—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Behavioral Risk Factor Surveillance System (BRFSS) is a CDC-sponsored system of cross-sectional telephone health surveys concerning individual health risk behaviors, health conditions, and preventive health practices that are associated with chronic diseases, infectious diseases, and injury. The BRFSS is administered annually by health departments in states, territories, and the District of Columbia (collectively referred to as states). An independent sample of respondents is drawn for each state. The system is designed to produce information that is specific to the public health needs of each participating jurisdiction, and for many is the only source of health risk data amenable to their uses. Although national estimates of some health risk behaviors are available, the methods used to produce national estimates do not typically produce the type of detailed information needed to plan and implement public health programs; moreover, national estimates provide only limited insight into regional or state-specific variability in health status and risk factors. Over time the BRFSS has developed into an important data collection system that federal agencies rely on for state and local health information and to track national health objectives such as Healthy People. Through the BRFSS partnership, CDC has established standard protocols for BRFSS data collection which all states are

encouraged to adopt. These standards allow for state-to-state data comparisons as well as comparisons over time.

The BRFSS questionnaire is based on modular design principles to accommodate a variety of state-specific needs within a common framework. All participating states are required to administer a standardized core questionnaire which provides a set of shared health indicators for all BRFSS partners. The BRFSS core questionnaire consists of fixed core, rotating core, and emerging core questions. Fixed core questions are asked every year. Rotating core questions cycle on and off the core questionnaire during even or odd years, depending on the question. Emerging core questions are included in the core questionnaire as needed to collect data on urgent or emerging health topics such as influenza.

In addition, the BRFSS includes a series of optional modules on a variety of topics. In off-years when the rotating questions are not included in the core questionnaire, they are offered to states as an optional module. This framework allows each state to produce a customized BRFSS survey by appending selected optional modules to the core survey. States may select which, if any, optional modules to administer. As needed, CDC provides technical and methodological assistance to state BRFSS coordinators in the construction of their state-specific surveys.

The CDC and BRFSS partners produce a new set of state-specific BRFSS questionnaires each calendar year (*i.e.*, 2016 BRFSS questionnaires, 2017 BRFSS questionnaires, etc.). CDC submits an annual Change Request to OMB outlining updates to the BRFSS core survey and optional modules that have occurred since the previous year. Each state administers its BRFSS questionnaire throughout the calendar year. The BRFSS partnership thus results in a flexible, coordinated information collection system that is adaptive to national and state-specific needs.

The current estimated average burden for the core BRFSS interview is 15 minutes. For the optional modules, the estimated average burden per response varies by state and year, but is currently estimated at an additional 15 minutes. Finally, the BRFSS allows states to customize some portions of the questionnaire through the addition of state-added questions, which are neither reviewed nor approved by the CDC. State-added questions are not included in CDC's burden estimates.

CDC periodically updates the BRFSS core survey and optional modules as new modules or emerging core

questions are adopted. The purpose of this Revision request is to incorporate field testing into the approved information collection plan.

Field testing is the final check of changes in the questionnaire which have occurred in the preceding year. Field testing is conducted in a manner that mimics the full-scale project protocol, to the degree that is feasible. Field testing is the final means by which changes are made in data collection methods and data collection software is tested. Field tests are used to identify problems with instrument documentation or instructions, problems with conditional logic (*e.g.*, skip patterns), software errors or other implementation and usability issues. Field testing is conducted with all new modules, emerging core questions, sections which precede and/or follow any new or changed items and extant sections which are topically related. This testing is conducted to ensure that questions are not perceived as

redundant or overlapping. Extant sections of the questionnaire unrelated to new items do not require testing. The demographic questions on the core BRFSS survey are included on each field test.

Since the field test instrument changes annually, it will be submitted to OMB for approval as an additional Change Request prior to implementation. Field tests are typically conducted in a single state with appropriate computer-assisted telephone interview (CATI) capability. Individuals who participate in field testing are drawn from a different sample than individuals who participate in the BRFSS surveys.

The BRFSS was initially approved with annualized estimates of 1,643,227 responses and 255,915 burden hours inclusive of the core survey and optional modules. CDC is requesting an additional allocation of 900 responses and 9,210 burden hours to conduct the annual field test. After a brief screening

interview, approximately 400 respondents per year will be determined ineligible or will decline to participate. The estimated burden per response for these respondents is one minute. An additional 500 respondents will participate in both the screening interview and the actual field test. The estimated burden for these respondents is 45 minutes. In years when fewer new questions and/or changes are proposed to the BRFSS questionnaire, field testing will impose a lesser burden. The revised total annualized estimates are 1,644,127 responses and 265,125 burden hours.

Information collection is conducted primarily to support state and local health departments, which plan and evaluate public health programs at the state or sub-state level. Information collected through the BRFSS is also used by the federal government and other entities. Participation in the BRFSS and its field test is voluntary and there are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hr)	Total burden (in hr)
U.S. General Population	Landline Screener	440,486	1	1/60	7,341
	Cell Phone Screener	223,334	1	1/60	3,722
	Field Test Screener	400	1	1/60	7
Annual Survey Respondents(Adults >18 Years).	BRFSS Core Survey	494,650	1	15/60	123,662
	BRFSS Optional Modules	484,757	1	15/60	121,189
Field Test Respondents(Adults >18 Years)	Field Test Survey	500	1	45/60	375
	Total				256,296

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-16-16JO; Docket No. CDC-2016-0005]

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 proposed information collect project entitled "The Pregnancy Risk Assessment Surveillance System".

DATES: Written comments must be received on or before March 21, 2016.

ADDRESSES: You may submit comments, identified by Docket No. CDC-2016-0005 by any of the following methods:

Federal eRulemaking Portal:
Regulation.gov. Follow the instructions for submitting comments.

Mail: Leroy A. Richardson,
Information Collection Review Office,
Centers for Disease Control and

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MS-D74, Atlanta, Georgia 30329.

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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.