

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

Survey of Sexually Transmitted Disease (STD) Provider Practices in the United States—NEW—National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Each year, 19.7 million sexually transmitted diseases (STDs) occur in the U.S., half of which strike youth 15–24 years of age. The public health burden of STDs is compounded by their economic impact. In 2010, an estimated \$15.6 billion in direct medical costs were attributed to STDs. Undiagnosed and untreated STDs can lead to serious long-term health consequences, especially for adolescent girls and young adult women. For example, every year, about 24,000 young women

become infertile as a result of undiagnosed and untreated STDs.

There is no national survey that collects detailed information on the STD practices of physicians. The STD Provider Survey will collect much needed data from U.S. health care providers in five specialties: Primary care (including internal medicine), general or family practice, obstetrics/gynecology, emergency medicine, and pediatrics. Knowledge of provider practices relative to guidelines and state-level laws and policies will provide information useful to stakeholders at all levels regarding the delivery of STD preventive services and treatment by health care providers in the U.S. As providers are one of the few professionals who have face-to-face contact with persons infected with STDs, they are also a potential intervention point for attempts to reduce re-infection and halt the further transmission of STDs.

The purpose of this survey is to conduct a nationally representative survey of physicians in five specialties: Primary care (including internal medicine), general or family practice, obstetrics/gynecology, emergency medicine, and pediatrics. Our sample size of physicians will allow for

national estimates and comparisons among these five specialties. Additionally, the survey will provide national estimates for comparisons between providers in the public and private sectors. Information collected will also be used to determine STD prevention activities needed by type of providers (by specialty or public/private) based on findings related to screening and treatment practices for STDs including EPT.

The survey contains sections on the physician's specialty areas, primary practice setting, primacy practice policies, patient demographics, STD testing and diagnosis, STD care and treatment, and respondent demographics.

In an effort to better understand policies and practices for STD care delivery among medical providers, the surveys will be sent to a random sample of 5,000 U.S. physicians across several specialties using the American Medical Association Master file. Using a multimode design (mail and web), multiple reminders will be sent to non-responders in order to reach the target of 3,500 completed surveys. The total burden hours are 1,342. There is no cost 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 hours)
Physicians responding via Mail	STD Provider Survey	2,625	1	20/60
Physicians responding via Web	STD Provider Survey	875	1	32/60

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-17-17WE; Docket No. CDC-2017-0025]

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 a proposed information collection project titled "Knowledge, Attitudes, and Practices related to a Domestic Readiness Initiative on Zika Virus Disease." This project consists of telephone interviews with participants in Puerto Rico and the domestic U.S.

DATES: Written comments must be received on or before May 26, 2017.

ADDRESSES: You may submit comments, identified by Docket No. CDC-2017-0025 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

Knowledge, Attitudes, and Practices related to a Domestic Readiness Initiative on Zika Virus Disease—New—Office of the Associate Director for Communication (OADC), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Since late 2015, Zika has rapidly spread through Puerto Rico. As of November 2016, there have been 35,136 confirmed cases of Zika in Puerto Rico, with 2,797 cases among pregnant women and 67 cases of Guillain-Barré caused by Zika. In the continental United States, there have been 4,432 travel-associated cases of Zika and 185 locally-acquired Zika cases in Florida and Texas. Due to the urgent nature of this public health emergency, CDC is implementing a Zika prevention communication and education initiative in the continental United States and Puerto Rico.

CDC intends to request approval from the Office of Management and Budget (OMB) to conduct an assessment of a domestic U.S. and Puerto Rico-based communication and education initiative aimed at encouraging at-risk populations to prepare and protect themselves and their families from Zika virus infection. As part of the mission of CDC's Domestic Readiness Initiative on the Zika Virus Disease, CDC will assess the following communication and education objectives: (1) Determine the reach and saturation of the initiative's messages in Puerto Rico and the domestic U.S.; (2) measure the extent to which messages were communicated clearly across multiple channels to advance knowledge and counter misinformation; and (3) monitor individual and community-level awareness, attitudes and likelihood to follow recommended behaviors.

This data collection is related to Zika prevention efforts that have been and will be implemented in Puerto Rico and the domestic U.S. Specifically, CDC needs this assessment to ensure that Zika prevention campaigns effectively reach target audiences to educate individuals regarding Zika prevention behaviors. Ongoing evaluation is an important part of this program because it can inform awareness of campaign activities, how people perceive Zika as a health risk, and assess their uptake of recommended health behaviors after the campaign has been implemented.

These interviews can help articulate motivations for and against engaging in Zika prevention behaviors that are critical for preventing Zika-associated birth defects and morbidities.

Implementing changes based on results from this assessment is expected to facilitate program improvement and ensure the most efficient allocation of resources for this public health emergency.

CDC will launch a new Zika Virus Disease Domestic Readiness Initiative in the continental U.S. and Puerto Rico. The goal of this project is to determine knowledge, attitudes, and practices related to this initiative. CDC will use the findings to improve planning, implementation, refinements, and demonstrate outcomes of a Zika Domestic Readiness Initiative communication and education effort. CDC will also use the information to make recommendations for improving communication and education regarding the prevention and spread of the Zika virus. CDC will develop presentations, reports, and manuscripts to document the communication effort and provide the lessons learned to inform future and similar communication efforts.

The plan is to conduct 2,400 interviews 12 months post-launch of the campaign to assess long term outcomes of the initiative. CDC will conduct telephone interviews with a mix of closed-ended and open-ended questions with individuals domestically in the U.S. and in Puerto Rico. The purpose of this assessment is to assess core components of CDC's Zika response in communicating prevention behaviors and risk messages to the public about vector control services.

The following factors will be assessed:

- Knowledge about Zika virus and related prevention behaviors
- Self-efficacy in engaging in Zika prevention behaviors
- Engagement in Zika prevention behaviors (e.g., protective clothing use, condom use, and standing water removal)
- Risk perceptions of Zika

Researchers will analyze the data, and generate a report for leaders of the response to offer insights on the delivery of the communication campaign.

Results of this project will have limited generalizability. However, results of this evaluation should provide information that can be used to enhance and revise the existing program as well as offer lessons learned to inform infectious disease control programs that use education materials.

Authorizing legislation comes from Section 301 of the Public Health Service Act (42 U.S.C. 241). There is no cost to respondents other than their time to participate.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
U.S. Domestic Adults	Zika Readiness Initiative Survey	1,800	1	14/60	420
Puerto Rico Adults	Zika Readiness Initiative Survey	600	1	14/60	140
Total	2,400	560

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 Medicare & Medicaid Services

[Document Identifier: CMS-10120]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare &
Medicaid Services, HHS.

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected; and the use of automated collection techniques or other forms of information technology to

minimize the information collection burden.

DATES: Comments on the collection(s) of information must be received by the OMB desk officer by April 26, 2017.

ADDRESSES: When commenting on the proposed information collections, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be received by the OMB desk officer via one of the following transmissions: OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer, Fax Number: (202) 395-5806 OR, Email: OIRA_submission@omb.eop.gov.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' Web site address at Web site address at <https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html>.

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@cms.hhs.gov.

3. Call the Reports Clearance Office at (410) 786-1326.

FOR FURTHER INFORMATION CONTACT: William Parham at (410) 786-4669.

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. The term "collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires federal agencies

to publish a 30-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* 1932(a) State Plan Amendment Template, State Plan Requirements and Supporting Regulations; *Use:* Section 1932(a)(1)(A) of the Social Security Act (the Act) grants states the authority to enroll Medicaid beneficiaries on a mandatory basis into managed care entities and primary care case managers. Under this authority, a state can amend its Medicaid state plan to require certain categories of Medicaid beneficiaries to enroll in managed care entities without being out of compliance with section 1902 of the Act on state-wideness (42 CFR 431.50), freedom of choice (42 CFR 431.51) or comparability (42 CFR 440.230). The template may be used by states to modify their state plans if they choose to implement the provisions of section 1932(a)(1)(A); *Form Number:* CMS-10120 (OMB control number: 0938-0933); *Frequency:* Once and occasionally; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 56; *Total Annual Responses:* 12; *Total Annual Hours:* 70. (For policy questions regarding this collection contact Debbie Anderson at 410-786-5545.)

Dated: March 22, 2017.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

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