DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–17–17ABU]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled *Zika Reproductive Health Call-Back Survey (ZRHCS), Puerto Rico, 2017* to the Office of Management and Budget (OMB) for review and approval. CDC previously published a ''Proposed Data Collection Submitted for Public Comment and Recommendations'' notice on April 27, 2017 to obtain comments from the public and affected agencies. CDC received one general comment related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

(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.* Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street NW., Washington, DC 20503 or by fax to (202) 395–5806. Provide written comments within 30 days of notice publication.

Proposed Project

Zika Reproductive Health Call-Back Survey (ZRHCS), Puerto Rico, 2017—New—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

In May 2015, the World Health Organization reported the first local mosquito born transmission of Zika virus in the Western Hemisphere, with autochthonous cases identified in Brazil. In response to the Zika virus outbreak, and evidence that Zika virus infection during pregnancy is a cause microcephaly and other adverse pregnancy and infant outcomes, CDC activated its Emergency Operations Center to its highest level on February 8, 2016 and continues to engage in Zika virus operations.

To date, Puerto Rico has reported the highest number of Zika virus cases of any area within the United States, with the Puerto Rico Department of Health (PRDH) reporting more than 40,000 cases of Zika virus infection, including 3,757 cases in pregnant women. Given the adverse pregnancy and birth outcomes associated with Zika virus infection during pregnancy and the current lack of a vaccine, it is important for women who are at risk of becoming pregnant unintentionally, or who are planning a pregnancy, to be knowledgeable about the potential outcomes of Zika virus infection. In addition, it is important for them to practice effective pregnancy prevention behaviors when they do not desire pregnancy and to prevent mosquito-borne and sexual transmission of Zika virus.

This is a request for a new information collection. CDC requests one additional year of clearance to continue the Emergency information collection, ''Emergency Zika Package: Zika Reproductive Health Survey, Puerto Rico, 2017,'' approved by the Office of Management and Budget (OMB) in July 2017 (OMB Control Number 0920–1188).

The objective of this assessment is to collect current information on various aspects of Zika knowledge and prevention behaviors from a representative sample of adult women in Puerto Rico. Information will be collected on the following topics: (1) Knowledge of and adherence to mosquito prevention strategies, and (2) use of condoms to minimize the risk of sexual transmission of Zika, and (3) behaviors practiced by women who wish to avoid or delay pregnancies that help them prevent unintended pregnancies that might otherwise be affected by Zika. CDC will rapidly summarize and analyze the information collected for the Puerto Rico Department of Health to determine the need for further refinements in educational messaging and allocation of resources, as established during the first season of the Zika outbreak. There is no cost to respondents other than the time to participate. The total estimated annual burden hours are 117.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Women aged 18–49 years who completed the main PR–BRFSS survey.	Recruitment text	645	1	1/60
Women aged 18–49 years who completed the main PR–BRFSS survey agree to participate in the call-back survey.	Call-back Survey and Consent.	581	1	10/60
PR–BRFSS Coordinators	Data Submission Layout	1	3	3

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–17–1122; Docket No. CDC–2017–0070]

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 reinstatement of the data collection project titled ''Congenital Heart Surveillance to Recognize Outcomes, Needs and well-being (CHSTRONG).'' CDC collects CHSTRONG data to provide public health question insight, aid in the development of services, and inform for the proper allocation of resources to improve long-term health and wellbeing.

DATES: Written comments must be received on or before November 20, 2017.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2017–0070 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 Leroy A. Richardson, 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

Congenital Heart Survey To Recognize Outcomes, Needs, and well-being (CH STRONG) (OMB Control Number: 0920–1122, Expiration 07/31/2017)—Reinstatement with change—National Center on Birth Defects and Developmental Disabilities (NCBDDD), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Congenital heart defects (CHDs) are the most common type of structural birth defects, affecting approximately 1 in 110 live-born children. In prior decades, many CHDs were considered fatal during infancy or childhood, but with tremendous advances in pediatric cardiology and cardiac surgery, at least 85% of patients now survive to adulthood and there are approximately 1.5 million adults with CHD living in the United States.

With vast declines in mortality from pediatric heart disease over the past 30 years, it is vital to evaluate long-term outcomes and quality of life issues for adults with CHD. However, U.S. data on long-term outcomes, quality of life issues, and comorbidities of adults born with CHD are lacking. U.S. data is needed to provide insight into the public health questions that remain for this population and to develop services and allocate resources to improve long-term health and wellbeing.

The initial request for this project was one year, but there was a delay in recruitment that results in a change in the recruitment process. Therefore, an additional 24 months is being requested. The three sites decided to conduct more intensive and time-consuming tracking and tracing to identify more accurate contact information for all eligible individuals. In addition to more intensive tracking and tracing, the sites decided to send recruitment materials in batches rather than all at once. This ensured that problems with the recruitment process were caught immediately and could be modified in subsequent rounds of recruitment. Due to these delays and changes in recruitment process, CH STRONG data collection is expected to last an additional 24 months and conclude two years after receiving an extension from OMB.