

includes health status and medical conditions, health care services, health behaviors, and sociodemographic characteristics. In addition, permission for collecting hospital discharge data, including diagnoses at discharge and procedures performed during hospitalization will be obtained during the interview.

Following the interview, a health examination will be conducted as part of the home visit. The respondent's weight, waist circumference, and sitting blood pressure will be measured, and a monofilament assessment may be conducted for neuropathy. In addition,

blood and urine will be collected. Examples of laboratory tests planned include hemoglobin A1c from the blood specimen, and albumin and creatinine from the urine collection. This proposed project will assess the feasibility of conducting these tests and procedures in the home examination setting.

A proxy interview will be conducted via telephone for sampled participants who died prior to the re-contact. Information on medical conditions and overnight hospital stays since baseline will be collected.

Although permission will be sought from all field feasibility test

participants, hospitalization records will be obtained only for 120 participants annually (240 participants over the 2-year period) to evaluate the record retrieval protocol for the study cohort among different medical facilities. An average of 3 hospital stays per person is anticipated among this cohort, therefore, an estimated 360 requests (120 persons  $\times$  3 stays) will be made annually. The estimated burden for hospital record provider is 20 minutes per record.

There is no cost 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 hours)	Total burden hours
2007–2014 NHANES examinees .....	Field feasibility test initial contact and appointment scheduling form.	400	1	20/60	133
2007–2014 NHANES examinees .....	Field feasibility test home visit .....	356	1	1	356
2007–2014 NHANES examinees .....	Field feasibility test home urine collection.	356	1	10/60	59
Proxy of deceased 2007–2014 NHANES examinees.	Field feasibility test deceased proxy interview.	44	1	20/60	15
Hospital record providers .....	Field feasibility test hospital records form.	360	1	20/60	120
Adult volunteers (non-field feasibility test participants).	Targeted methodological studies .....	375	1	1	375
Total .....	.....	.....	.....	.....	1,058

#### 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

[30 Day–16–0987]

#### 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](mailto: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

Qualitative Information Collection on Emerging Diseases among the Foreign-born in the U.S. (OMB Control No. 0920–0987, Expires 09/30/2016)—Extension—Division of Global Migration and Quarantine, National Center for Emerging Zoonotic and Infectious Diseases, Centers for Disease Control and Prevention (CDC).

#### Background and Brief Description

The Centers for Disease Control and Prevention (CDC), National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Division of Global Migration and Quarantine (DGMQ), requests approval for an extension of the current generic information collection Qualitative Information Collection on Emerging Diseases among the Foreign-born in the U.S. (OMB Control Number 0920–0987, expiration date 9/30/2016).

This qualitative data collection is needed by DGMQ because foreign-born individuals are considered hard-to-reach populations and are often missed by routine information collection systems in the United States. As a consequence, limited information is available about the health status, knowledge, attitudes, health beliefs and practices related to communicable diseases and other emerging health issues (e.g., tuberculosis, parasitic diseases, lead poisoning, and mental health issues) among foreign-born populations in the United States. Foreign-born populations are very diverse in terms of countries of origin, socio-demographic, cultural and linguistic characteristics and geographic destinations in the U.S. Data is especially limited at the local level.

The purpose of the extension is to continue efforts to improve the agency's understanding of the health status, risk factors for disease, and other health outcomes among foreign-born individuals in the United States. Numerous types of data will be collected under the auspices of this generic information collection. These include, but are not limited to, knowledge, attitudes, beliefs, behavioral intentions, practices, behaviors, skills, self-efficacy, and health information needs and sources.

For example, CDC recently used this generic to collect feedback on Mexican-born audience's preferences for messaging and communication about mosquito-borne diseases to develop effective prevention campaigns as these diseases—especially Zika—pose an

increasing threat to global health security.

Under the terms of this generic, CDC will employ focus groups and key informant interviews to collect information. Depending on the specific purpose, the information collection may be conducted either in-person, by telephone, on paper, or online. For each generic information collection, CDC will submit to OMB the project summary and information collection tools.

#### *Estimated Annualized Burden Hours*

This requests entails a total of 1,025 respondents and 825 burden hours annually. The respondents to these information collections are foreign born individuals in the United States. There is no cost to respondents other than the time required to provide the information requested.

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Foreign-born from specific country of birth in the United States.	Screeners for focus groups (assuming 2 screenings for each recruited participant in focus groups) (300X2 = 600).	600	1	10/60
Foreign-born from specific country of birth in the United States.	Focus Groups (Approximately 30 focus groups/year and 10 participants per focus group).	300	1	2
Foreign-born community leaders and staff from organizations serving those communities.	Key informant interviews (Approximately 125 interviews/year).	125	1	1

**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-16-0950; Docket No. CDC-2016-0044]

### 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 revision of the National Health and Nutrition Examination Survey (NHANES). NHANES programs produce descriptive statistics which measure the health and nutrition status of the general population.

**DATES:** Written comments must be received on or before July 22, 2016.

**ADDRESSES:** You may submit comments, identified by Docket No. CDC-2015-0044 by any of the following methods:

- *Federal eRulemaking Portal:* [Regulation.gov](http://www.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*.

**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](mailto: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