As the outbreak evolves, interviews with pregnant women in Puerto Rico 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 has occurred with the previous information collection and is expected to facilitate program improvement and ensure the most efficient allocation of resources for this public health emergency. Understanding risk and protective factors related to interventions and behaviors of pregnant women can help to establish priorities.

There are no costs to the respondents other than their time. The total number of estimated annual burden hours is 500.

**ESTIMATED ANNUALIZED BURDEN HOURS**

<table>
<thead>
<tr>
<th>Type of respondents</th>
<th>Form name</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden per response (hrs.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pregnant WIC participant</td>
<td>Initial Telephone Interview</td>
<td>1,200</td>
<td>1</td>
<td>20/60</td>
</tr>
<tr>
<td></td>
<td>Follow-up Telephone Interview</td>
<td>600</td>
<td>1</td>
<td>10/60</td>
</tr>
</tbody>
</table>

**Department of Health and Human Services**

**Centers for Disease Control and Prevention**

**[30Day–17–16AXC]**

**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**

Assessing Safety and Health Hazards to Workers in Oil and Gas Extraction: A Survey—New—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

**Background and Brief Description**

The mission of the National Institute for Occupational Safety and Health (NIOSH) is to promote safety and health at work for all people through research and prevention. The Occupational Safety and Health Act, 91 (section 20[a] [1]), authorizes NIOSH to conduct research to advance the health and safety of workers. NIOSH is proposing a two year study to conduct a survey questionnaire of 500 land-based oil and gas (O&G) extraction workers in 5 U.S. states (Texas, North Dakota, Colorado, Oklahoma and a state in the Appalachian Basin) to examine safety and health issues and concerns of this workforce. Workers who drive as a part of their work duties will be asked to complete an additional set of questions about their driving environment and health issues and concerns of this workforce.

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.

[FR Doc. 2017–06868 Filed 4–5–17; 8:45 am]
who drive a company vehicle will also be asked to complete “Module 4: Motor Vehicle.” An estimated 75% of the workers will complete the driving portion of the survey (187 workers). This module will take approximately 5 additional minutes to complete for those using the tablet (approximately 168 workers per year) as well as 5 minutes for those completing the hardcopy version (19 workers per year).

Comments submitted in response to this notice will be reviewed and addressed prior to OMB application submission. The total estimated burden hours are 151.

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**ESTIMATED ANNUALIZED BURDEN HOURS**

<table>
<thead>
<tr>
<th>Type of respondents</th>
<th>Form name</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden per response (in hours)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Screening of Workers</td>
<td>Non Respondent Questionnaire</td>
<td>313</td>
<td>1</td>
<td>5/60</td>
</tr>
<tr>
<td>O&amp;G Extraction Workers</td>
<td>Tablet Version, Modules 2: General, Module 3: Well Site Work, and, Module 5: Closing Questions.</td>
<td>63</td>
<td>1</td>
<td>5/60</td>
</tr>
<tr>
<td>O&amp;G Extraction Workers</td>
<td>Hardcover, Version, Modules 2: General, Module 3: Well Site Work, and, Module 5: Closing Questions.</td>
<td>225</td>
<td>1</td>
<td>25/60</td>
</tr>
<tr>
<td>O&amp;G Extraction Workers who drive at work</td>
<td>Tablet Version, Module 4: Motor Vehicle</td>
<td>25</td>
<td>1</td>
<td>25/60</td>
</tr>
<tr>
<td>O&amp;G Extraction Workers who drive at work</td>
<td>Hardcover Version, Module 4: Motor Vehicle</td>
<td>168</td>
<td>1</td>
<td>5/60</td>
</tr>
</tbody>
</table>

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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.

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 Zika virus persistence in body fluids of patients with Zika virus infection in Puerto Rico (ZIPER Study) to fill gaps in the scientific knowledge base for ZIKV regarding the persistence and transmissibility of ZIKV in body fluids. This information assist ongoing public health response activities, as well as advance the scientific understanding of ZIKV illness and transmission.

DATES: Written comments must be received on or before June 5, 2017.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2017–0030 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, install and utilize technology and systems for the purpose of collecting, validating and verifying