Jeffrey M. Zirger,

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Centers for Disease Control and Prevention
[30Day–18–18TH]

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 “Assessment of a Preventive Service Program in the Context of a Zika Virus Outbreak in Puerto Rico” 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 March 30, 2018 to obtain comments from the public and affected agencies. CDC received one non-substantive comment 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

Assessment of a Preventive Service Program in the Context of a Zika Virus Outbreak in Puerto Rico—New—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Puerto Rico has reported the highest number of Zika virus infections in the United States, including infections in pregnant women. Zika virus infection during pregnancy has been identified as a cause of microcephaly and other severe brain abnormalities, and has been linked to other problems such as miscarriage, stillbirth, defects of the eye, hearing deficits, limb abnormalities, and impaired growth. One strategy to prevent these devastating outcomes is to prevent unintended pregnancy among women at risk of Zika virus infection.

To this end, an initiative was launched in April 2016 to train physicians at clinics across Puerto Rico to provide patient-centered services to women who chose to delay or avoid pregnancy during the Zika virus outbreak.

As part of the public health response to the Zika virus outbreak, CDC seeks to assess approaches to mitigating the effects of Zika virus infection and determine which approaches have utility. Previous assessment of the prevention program indicated high satisfaction of patients with program services. The specific objectives of this data collection are to assess (1) prevention strategy adherence among patients at approximately 18 months after receipt of program services; and (2) prevention strategy adherence, patient satisfaction, and unmet need for services among participants at approximately 30 months after receipt of program services. The practical utility of the information to be collected as part of this project is to assess services delivered to women in Puerto Rico, monitor outcomes of interest, and determine potential for replication/adaptation in other jurisdictions similarly affected by the Zika virus or during other emergency responses. For the information collection, CDC plans to conduct online surveys with 1,920 patients approximately 18 months after receiving program services and 1,760 patients approximately 30 months after receiving program services. The number of patients surveyed is based on an initial sample of 3,200 patients invited to participate, anticipating a 60% response rate at 18 months and a 55% response rate at 30 months.

Participation in all data collection activities will be completely voluntary. OMB approval is requested for two years. Total Annualized Burden Hours are estimated to be 259, and there are no costs to respondents other than their time.
### DEPARTMENT OF HEALTH AND HUMAN SERVICES

**Centers for Disease Control and Prevention**

[**30Day–18–1091**]

**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 “Using Qualitative Methods to Understand Issues in HIV Prevention, Care and Treatment in the United States” 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 March 13, 2018 to obtain comments from the public and affected agencies. CDC received four comments 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

Using Qualitative Methods to Understand Issues in HIV Prevention, Care and Treatment in the United States (OMB No. 0920–1091; expires December 31, 2018)—Extension—National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention, Centers for Disease Control and Prevention (CDC).

### Background and Brief Description

The CDC’s National Center on HIV/AIDS, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), Division of HIV/AIDS Prevention (DHAP) seeks a three year extension for an existingGeneric information collection request (Generic ICR) entitled, “Using Qualitative Methods to Understand Issues in HIV Prevention, Care and Treatment in the United States” (OMB Number: 0920–1091). Specific studies conducted under this extended Generic ICR will be consistent with the national HIV prevention goals, the CDC Division of HIV/AIDS Prevention (DHAP) Strategic Plan, and DHAP’s High-impact HIV Prevention approach.

The purposes for each data collection study supported under this extended Generic ICR will be to understand specific barriers and facilitators to local HIV prevention, care and treatment in the United States and territories. For example, each study will seek to identify ways to improve programmatic activities along the continuum of HIV prevention, treatment and care for different populations residing in different geographic settings with greatest burden of HIV.

The target populations for studies included in this extended Generic ICR include, but are not limited to: Persons living with HIV who are in treatment; persons living with HIV who are out of treatment and who may or may not be seeking treatment at healthcare facilities; persons at high risk for HIV acquisition (HIV negative) and HIV transmission (HIV positive); persons from groups at high risk for HIV including gay, bisexual and other MSM, transgender persons, and injection and non-injection drug users; persons from racial and ethnic minorities; and healthcare providers or other professionals who provide HIV prevention, care and treatment services. Other populations may include individuals who provide non-HIV services or otherwise interact with persons living with HIV or persons at risk for HIV acquisition.

Studies will only provide local contextual information about the barriers and facilitators to HIV prevention, care, and treatment experienced by specific communities at risk for acquiring HIV infection, by HIV-positive persons across the HIV care continuum, and by organizations or individuals providing HIV prevention, care, treatment, and related support services.

Data collection methods used in any of the specific studies primarily will consist of rapid qualitative assessment methodologies, such as semi-structured and in-depth qualitative interviews, focus groups; direct observations; document reviews; and short structured surveys. Data will be analyzed using

<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>Patients aged 18 years or older</td>
<td>Online surveys (18-month follow-up)</td>
<td>960</td>
<td>1</td>
<td>7/60</td>
</tr>
<tr>
<td>Patients aged 18 years or older who completed 18 mo survey.</td>
<td>Online surveys (30-month follow-up)</td>
<td>660</td>
<td>1</td>
<td>10/60</td>
</tr>
<tr>
<td>Patients aged 18 years or older who did not complete 18 mo survey.</td>
<td>Online surveys (30-month follow-up)</td>
<td>220</td>
<td>1</td>
<td>10/60</td>
</tr>
</tbody>
</table>

**Jeffrey M. Zirger,**