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

DELTA FOCUS Program Evaluation (OMB No. 0920–0984)—Reinstatement with Change—National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC).

**Background and Brief Description**

Intimate Partner Violence (IPV) is a serious, preventable public health problem that affects millions of Americans and results in serious consequences for victims, families, and communities. IPV occurs between two people in a close relationship. The term “intimate partner” describes physical, sexual, or psychological harm by a current or former partner or spouse. IPV can impact health in many ways, including long-term health problems, emotional impacts, and links to negative health behaviors. IPV exists along a continuum from a single episode of violence to ongoing battering; many victims do not report IPV to police, friends, or family. In 2002, authorized by the Family Violence Prevention Services Act (FPVSA), CDC developed the Domestic Violence Prevention Enhancements and Leadership Through Alliances (DELTA) Program, with a focus on the primary prevention of IPV.

The purpose of the DELTA FOCUS program is to promote the prevention of IPV through the implementation and evaluation of strategies that create a foundation for the development of practice-based evidence. By emphasizing primary prevention, this program will support comprehensive and coordinated approaches to IPV prevention. On March 2, 2013, CDC awarded 10 cooperative agreements to state domestic violence coalitions (SDVCs).

Each SDVC is required to identify and fund one to two well-organized, broad-based, active local organizations (referred to as coordinated community responses or CCRs) that are already engaging in, or are at capacity to engage in, IPV primary prevention strategies affecting the structural determinants of health at the societal and/or community levels of the SEM. SDVCs must facilitate and support local-level implementation and hire empowerment evaluators (EEs) to support the evaluation of IPV prevention strategies by the CCRs. SDVCs must also implement and with their empowerment evaluators, evaluate state-level IPV prevention strategies.

The CDC seeks OMB approval for three years to collect program evaluation data. Information will be collected from awardees funded under FOA–CE13–1302, the DELTA FOCUS (Domestic Violence Prevention Enhancement and Leadership Through Alliances, Focusing on Outcomes for Communities United with States) cooperative agreement program. The information will be used to guide program improvements by CDC in the national DELTA FOCUS program implementation and program improvements by SDVCs in implementation of the program within their state. Not collecting this data could result in inappropriate implementation, resulting in ineffective use of tax payer resources. Thus, this data collection is an essential program evaluation activity and the results will not be generalizable to the universe of study. The estimated annual burden hours are 59. There is no cost to respondents other than their time.

<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 hrs.)</th>
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<tbody>
<tr>
<td>Director of Nursing, Registered Nurse, Infection Control and Prevention Officer. Registered Nurse</td>
<td>Healthcare Facility Assessment. Residents by Location Form</td>
<td>200</td>
<td>1</td>
<td>45/60</td>
</tr>
<tr>
<td>Licensed Practical or Licensed Vocational Nurses</td>
<td></td>
<td>200</td>
<td>38</td>
<td>20/60</td>
</tr>
</tbody>
</table>
### ESTIMATED ANNUALIZED BURDEN HOURS

<table>
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<tr>
<th>Type of respondent</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>DELTA FOCUS Awardees (SDVC executive directors, SDVC project coordinators, SDVC empowerment evaluators, and SDVC-funded CCR project coordinators).</td>
<td>DELTA FOCUS Survey</td>
<td>59</td>
<td>1</td>
<td>1</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.

[FR Doc. 2016–12706 Filed 5–27–16; 8:45 am]

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–855(A, B, I)]

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

**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 a 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 any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency’s functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) 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 June 30, 2016.

**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:


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:** Reports Clearance Office at (410) 786–1326.

**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: Extension of a currently approved collection; Title of Information Collection: Medicare Enrollment Application; Use: The primary function of the CMS–855 Medicare enrollment application is to gather information from a provider or supplier that tells us who it is, whether it meets certain qualifications to be a health care provider or supplier, where it practices or renders its services, the identity of the owners of the enrolling entity, and other information necessary to establish correct claims payments. No comments were received during the 60-day comment period (April 1, 2016 (81 FR 18855)). Form Number: CMS–855(A, B, I) (OMB control number: 0938–0685); Frequency: Annually; Affected Public: Private Sector; Business or other for-profit and not-for-profit institutions; Number of Respondents: 1,735,800; Total Annual Responses: 86,480; Total Annual Hours: 290,193. (For policy questions regarding this collection contact Kimberly McPhillips at 410–786–5374.)


**William N. Parham, III,**

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

[FR Doc. 2016–12694 Filed 5–27–16; 8:45 am]

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

Food and Drug Administration

[Docket No. FDA–2016–N–0001]

Advisory Committee; Allergenic Products Advisory Committee, Renewal

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice; renewal of advisory committee.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing the renewal of the Allergenic Products Advisory Committee by the Commissioner of Food and Drugs (the Commissioner). The Commissioner has determined that it is in the public interest to renew the Allergenic Products Advisory Committee for an additional 2 years beyond the charter