### DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### Centers for Disease Control and Prevention

**[30Day-16–16TM]**

#### 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

**Prevalence Survey of Healthcare-Associated Infections and Antimicrobial Use in U.S. Nursing Homes—New—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).**

**Background and Brief Description**

Preventing healthcare-associated infections (HAI) and encouraging appropriate use of antimicrobials are priorities of both the U.S. Department of Health and Human Services and the Centers for Disease Control and Prevention. The burden and epidemiology of HAIs and antimicrobial use in U.S. nursing homes is currently unknown. Understanding the scope and magnitude of all types of HAIs in patient populations across the spectrum of U.S. healthcare facilities is essential to the development of effective prevention and control strategies and policies.

HAI prevalence and antimicrobial use estimates can be obtained through prevalence surveys in which data are collected in healthcare facilities during a short, specified time period. Essential steps in reducing the occurrence of HAIs and the prevalence of resistant pathogens include estimating the burden, types, and causative organisms of HAIs; assessing the nature and extent of antimicrobial use in U.S. healthcare facilities; and assessing the nature and extent of antimicrobial use.

Prevalence surveys, in which data are collected in healthcare facilities during a short, specified time period represent an efficient and cost-effective alternative to prospective studies of HAI and antimicrobial use incidence. Given the absence of existing HAI and antimicrobial use data collection mechanisms for nursing homes, prevalence surveys represent a robust method for obtaining the surveillance data required to identify HAIs and antibiotic use practices that should be targeted for more intensive surveillance and to guide and evaluate prevention efforts.

The methods for the data collection are based on those used in CDC hospital prevalence surveys and informed by a CDC pilot survey conducted in nine U.S. nursing homes. The survey will be performed by the CDC through the Emerging Infections Program (EIP), a collaboration with CDC and 10 state health departments with experience in HAI surveillance and data collection. Respondents are nursing homes certified by the Centers for Medicare & Medicaid Services in EIP states. Nursing homes will be randomly selected for participation. The EIP will recruit 20 nursing homes in each of the 10 EIP sites. Nursing home participation is voluntary.

OMB approval is requested for three years. Participation is voluntary and there are no costs to respondents other than their time. The total estimated annual burden hours are 5,217.

#### ESTIMATED ANNUALIZED BURDEN HOURS—Continued

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<th>Type of respondents</th>
<th>Form name</th>
<th>Number of respondents</th>
<th>Number of responses per Respondent</th>
<th>Avg. burden per response (in hrs.)</th>
<th>Total burden (in hrs.)</th>
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<td>Total</td>
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<td>5,110,716</td>
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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.

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**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 (FVPSA), 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 two 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.