

“Varicella (Chickenpox) Vaccine: What You Need to Know,” publication date August 6, 2021.

“Your Child’s First Vaccines: What You Need to Know,” publication date October 15, 2021.

With publication of this notice, by March 31, 2022, all healthcare providers must discontinue use of the previous editions and provide copies of these updated vaccine information materials prior to immunization in conformance with CDC Instructions for Use of Vaccine Information Statements dated October 15, 2021.

Dated: December 20, 2021.

Angela K. Oliver,

Executive Secretary, Centers for Disease Control and Prevention.

[FR Doc. 2021-27929 Filed 12-23-21; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-22-0852]

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 Prevalence Survey of Healthcare-Associated Infections and Antimicrobial Use in U.S. Acute Care Hospitals 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 August 13, 2021 to obtain comments from the public and affected agencies. CDC received one non-substantive comment 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. Comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function. 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

Prevalence Survey of Healthcare-Associated Infections and Antimicrobial Use in U.S. Acute Care Hospitals (OMB Control No. 0920-0852, Exp. 10/31/2022)—Extension—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Preventing healthcare-associated infections (HAIs) and improving antimicrobial use (AU) are CDC and national priorities. An essential step in reducing the occurrence of HAIs is to accurately estimate the burden of these infections in U.S. acute care hospitals and to describe the types of HAIs and causative pathogens. Periodic assessments of the magnitude and types of HAIs and AU occurring in all patient populations within acute care hospitals are needed to inform decisions by policy makers and hospital infection control personnel (ICP) regarding appropriate targets and strategies for HAI prevention and antimicrobial stewardship.

Since 2009, CDC has conducted four prevalence surveys (*i.e.*, pilot survey in 2009, limited-scale survey in 2010, and two full-scale surveys in 2011 and 2015)

in partnership with the CDC’s Emerging Infections Program (EIP) sites. Findings from the most recent survey showed a reduction in the percentage of patients with healthcare-associated infections compared with 2011. We granted approval from OMB to conduct the fifth survey in 2020, but due to the COVID-19 pandemic the survey was postponed to 2023.

Minor adjustments to data collection instruments since the previous 2019 OMB approval have been made. These adjustments were made to enhance future analyses and utility of the survey data. These changes are non-substantive and are not expected to increase the public reporting burden. An extension of the prevalence survey’s existing OMB approval is sought to allow a repeat HAI and AU Prevalence Survey to be performed in 2023. A repeat survey will allow assessment of changes in HAI and AU prevalence, pathogen distribution, and quality of antimicrobial prescribing. These data will also allow CDC and its partners to continue to monitor HAI and AU trends, to measure progress in meeting national targets, and to further refine prevention strategies.

In the 2023 survey, data collection will occur within acute care general hospitals of varying size in each of the 10 EIP sites (*i.e.*, CA, CO, CT, GA, MD, MN, NM, NY, OR, & TN). Infection Control Personnel in participating hospitals may assist EIP site personnel in collecting demographic and limited clinical data from the electronic or paper-based medical records of a sample of randomly selected patients on a single day in 2023. Patients will not be interviewed, and no direct interaction with patients will occur. Hospital and patient-level data will be collected using unique identification codes. EIP site personnel will submit hospital and patient-level data to CDC using a secure data management system.

Based on experiences from previous surveys, the time required to complete the Healthcare Facility Assessment Form (HFA) and Patient Information Form (PIF) is estimated to be 45 and 17 minutes, respectively. To conduct the full-scale survey in a three-year approval period, 100 hospital respondents will complete the HFA once, and the PIF on average 63 times per year. The total estimated annualized public burden is 1,860 hours, which represents no change from the 2019 OMB approval.

To assess changes in HAIs and AU over time, EIP sites will seek participation from the same hospitals that participated in prior surveys. These hospitals were originally selected for participation using a stratified random

sampling scheme based on the number of staffed acute care beds (*i.e.*, small: <150 staffed beds; medium: 151–399 staffed beds; large: >400 staffed beds). Each site will also have the option to recruit additional hospitals for a total of

up to 30 in each site. As in previous surveys, hospital participation will remain voluntary. Within each participating hospital, EIP site personnel will establish patient sample size targets based on the number of

staffed acute care beds (*e.g.*, up to 75 patients in small hospitals, 75 patients in medium hospitals, and 100 patients in large hospitals).

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Hospital Staff (Infection Preventionist)	HFA	100	1	45/60
	PIF	100	63	17/60

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–22–0017]

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 “Application for Training” 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 July 26, 2021 to obtain comments from the public and affected agencies. CDC received one comment related to the previous notice, and provided a standard response. 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. Comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function. 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

Application for Training (OMB Control No. 0920–0017, Exp. 4/30/2022)—Revision—Center for Surveillance, Epidemiology, and Laboratory Services (CSELS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The mission of CDC’s Division of Scientific Education and Professional Development (DSEPD) is to support the development of a competent, sustainable, and empowered public health workforce. Professionals in public health, epidemiology, medicine,

economics, information science, veterinary medicine, nursing, public policy, and other related professions seek professional development opportunities (both accredited and nonaccredited) through two CDC learning management systems. These two learning management systems are Training and Continuing Education Online (TCEO) (for accredited courses) and CDC TRAIN (for nonaccredited courses developed by CDC programs, grantees, and other funded partners). These two systems allow for the public health workforce to broaden their knowledge and skills to improve the science and practice of public health for domestic and international impact. Both systems currently involve related, but separate, information collection tools and information technology platforms.

The CDC seeks approval to implement changes as follows:

1. In TCEO, two additional accreditation types will be added as options a learner can select, to allow for master certified health education specialists and physician assistants to earn continuing education. Additional text is added to clarify what is requested for the CPE (Continuing Pharmacy Education) ID number.

2. CDC TRAIN is added as a data collection platform. The addition of CDC TRAIN to this request also supports the eventual merger of the two learning systems, a process that is underway and described further below. Adding CDC TRAIN to this revision also would allow CDC programs to collect standardized post-course evaluation data for program improvement, similar to what is done currently in TCEO (see #3).

3. The two standard training evaluation tools in CDC TRAIN are added to evaluate a training’s effectiveness (learning transfer and quality training) as well as its promotion, delivery, and learner satisfaction at two time points