

necessity and utility of the proposed information collection for the proper performance of the Agency's function; (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.

1. Type of Information Collection

Request: Extension of a currently approved collection; *Title of Information Collection:* Organ Procurement Organization's (OPO's) Health Insurance Benefits Agreement and Supporting Regulations at 42 CFR 486.301–486.348; *Use:* The information provided on this form serves as a basis for continuing the agreements with CMS and the 580 OPOs for participation in the Medicare and Medicaid programs for reimbursement of service. *Form Number:* CMS–576A (OMB#: 0938–0512); *Frequency:* Reporting—Occasionally; *Affected Public:* Private Sector: Business or other for-profits and Not-for-profit institutions; *Number of Respondents:* 58; *Total Annual Responses:* 58; *Total Annual Hours:* 116. (For policy questions regarding this collection contact Michele Walton at 410–786–3353. For all other issues call 410–786–1326.)

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS Web Site address at <http://www.cms.hhs.gov/PaperworkReductionActof1995>, or E-mail your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@cms.hhs.gov, or call the Reports Clearance Office on (410) 786–1326.

To be assured consideration, comments and recommendations for the proposed information collections must be received by the OMB desk officer at the address below, no later than 5 p.m. on March 31, 2010.

OMB, Office of Information and Regulatory Affairs,
Attention: CMS Desk Officer,
Fax Number: (202) 395–6974,
E-mail:
OIRA_submission@omb.eop.gov.

Dated: February 23, 2010.

Michelle Shortt,

*Director, Regulations Development Group,
Office of Strategic Operations and Regulatory Affairs.*

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–10–10AD]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639–5960 or send an e-mail to omb@cdc.gov. Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

School Dismissal Monitoring System—New—National Center for Preparedness, Detection, and Control of Infectious Diseases (NCPDCID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

During the spring 2009 H1N1 outbreak, the U.S. Department of Education (ED) and the Centers for Disease Control and Prevention (CDC) received numerous daily requests about the overall number of school dismissals nationwide including the number of students and teachers impacted by the outbreak. Illness among school-aged students (K–12) in many states and cities resulted in at least 1351 school dismissals due to rapidly increasing absenteeism among students or staff that impacted at least 824,966 students and 53,217 teachers.

Although a system was put in place to track school closures in conjunction with the Department of Education (ED), no formal monitoring system was established, making it difficult to monitor reports of school dismissal and to gauge the impact of the outbreak.

CDC has recently issued guidance for school closure for the 2009–2010 school year. To address the need to monitor reports of school closure, CDC and ED have established a School Dismissal Monitoring System to report on novel influenza A (H1N1)-related school or school district dismissals in the United States. Although the School Dismissal Monitoring System is currently approved to collect data under OMB Control Number 0920–0008, Emergency Epidemic Investigations, CDC would like to continue the data collection long term. Thus, CDC is requesting a separate OMB Control Number for this data collection.

The purpose of the School Dismissal Monitoring System is to generate accurate, real-time, national summary data daily on the number of school dismissals and the number of students and teachers impacted by the school dismissals. CDC will use the summary data to fully understand how schools are responding to CDC community mitigation guidance among schools, students, household contacts and for overall awareness of the impact of influenza outbreaks on school systems and communities.

Respondents are schools, school districts, and local public health agencies. Respondents will use a common reporting form to submit data to CDC. The reporting form includes the following data elements: Name of school district; zip code of school district; date the school or school district was dismissed; and the date school or school district is projected to reopen. Optional data elements include: name of person submitting information; the organization/agency; phone number of the organization/agency; and e-mail address. There is no cost to respondents other than their time to complete the data collection. The total annualized burden for this information collection request is 42 hours.

ESTIMATE OF ANNUALIZED BURDEN HOURS

Respondent	Number of respondents	Responses per respondent	Average burden per respondent (in hours)
School, school district or public health department	500	1	5/60

Dated: February 24, 2010.

Maryam I. Daneshvar,

Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. 2010-4177 Filed 2-26-10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-10-10BT]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404-639-5960 or send comments to Maryam I. Daneshvar, CDC Acting Reports Clearance Officer, 1600 Clifton Road, MS-D74, Atlanta, GA 30333 or send an e-mail to omb@cdc.gov.

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; and (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. Written comments should be received within 60 days of this notice.

Proposed Project

National Quitline Data Warehouse — New—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description:

Despite the high level of public knowledge about the adverse effects of smoking, tobacco use remains the leading preventable cause of disease and

death in the United States. Tobacco use results in approximately 440,000 deaths annually, including approximately 38,000 deaths from secondhand smoke exposure. Adults who smoke contribute to \$92 billion annually in lost worker productivity, and die an average of 14 years earlier than nonsmokers. Although the prevalence of current smoking among adults decreased significantly since its peak in the 1960s, overall smoking prevalence among U.S. adults has remained virtually unchanged during the past five years. Large disparities in smoking prevalence continue to exist among members of racial/ethnic minority groups and individuals of low socioeconomic status.

The National Tobacco Control Program (NTCP) was established by CDC to help reduce tobacco-related disease, disability, and death. The NTCP's four goal areas are: (1) The prevention of initiation of tobacco use among young people, (2) the elimination of nonsmokers' exposure to secondhand smoke, (3) the promotion of quitting among adults and young people, and (4) the elimination of tobacco-related disparities. The NTCP has provided funding for State quitlines, which provide telephone-based tobacco cessation services—including individualized counseling and self-help material—to help tobacco users quit. Quitlines overcome many of the barriers to tobacco cessation classes and traditional clinics because they are free and available at the caller's convenience. Quitline services in all States can be accessed through a toll-free national portal number at 1-800-QUIT-NOW. According to CDC's Best Practices for Comprehensive Tobacco Control, approximately six to eight percent of tobacco users potentially can be reached successfully by quitlines; however, currently, only one to two percent of tobacco users contact quitlines.

All States collect intake information about quitline callers and the services provided to them, but have varied with respect to the schedule for follow-up with callers, the number of follow-up attempts per caller, and the collection of information related to follow-up. With leadership from the North American Quitline Consortium (NAQC) and other tobacco control organizations, the field has collaborated to develop a Minimum Data Set (MDS) consisting of a set of suggested intake questions that should be asked of all callers, and follow-up questions that should be asked of a

representative sample of callers who have both completed intake and received a quitline service.

CDC requests OMB approval to collect information for a National Quitline Data Warehouse (NDQW) based on a uniform follow-up protocol and standardized instruments adapted from the MDS. Respondents will be the 50 States, the District of Columbia, and Guam. Additional funding for the expansion of tobacco quitline services, standardization of the information collection, and transmission to the shared NQDW is provided under the *American Recovery and Reinvestment Act of 2009* (ARRA).

Intake information will be collected from approximately 60,833 callers per month over a 24-month period. Minimal information will be collected from callers who contact the Quitline on behalf of another person. The information collection will also include seven-month follow-up data from a random sample of approximately 3,400 callers per month across all States, beginning in month eight (*i.e.*, seven full months after the first intakes) and continuing through month 24. Finally, the Tobacco Control Manager for each ARRA awardee (State, district or territory) will be required to submit a quarterly report describing services provided. The quarterly report will be used to quantify improvements in the capacity of the quitlines to assist tobacco users over time and to evaluate the expenditure of Recovery Act dollars.

The NQDW will have significant implications for the development of policies and programs aimed at tobacco use cessation and reduction of tobacco use. The information to be collected in the NQDW will be used to determine the role quitlines are playing in promoting tobacco use cessation, measure the number of tobacco users being served by State Quitlines, determine reach of quitlines to high-risk populations (*e.g.*, racial and ethnic minorities and the medically underserved), measure the number using each State quitline who quit, determine whether some combinations of services contribute to higher quit rates than others, and improve the timeliness, access to, and quality of data collected by quitlines.

CDC requests OMB approval to collect information for a two-year period. All information will be collected electronically. There are no costs to respondents other than their time.