Type of response	Number of respondents	Responses per respondent	Total responses	Hours per response	Total hour burden
Client-level baseline interview	15,681 10,646 4,508 2,352 9,017	1 1 1 1	15,681 10,646 4,508 2,352 9,017	0.333 0.367 0.367 0.1 0.1	5,222 3,907 1,655 235 902
Client-level Subtotal	15,681		15,681		11,920
Infrastructure development, prevention, and mental health promotion quarterly record abstraction	942	4	3,768	4	15,072
Total	16,623				26,992

### ESTIMATE OF ANNUAL RESPONSE BURDEN

Written comments and recommendations concerning the proposed information collection should be sent by April 23, 2010 to: SAMHSA Desk Officer, Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, DC 20503; due to potential delays in OMB's receipt and processing of mail sent through the U.S. Postal Service, respondents are encouraged to submit comments by fax to: 202–395–6974.

Dated: March 17, 2010.

### Elaine Parry,

Director, Office of Program Services.
[FR Doc. 2010–6457 Filed 3–23–10; 8:45 am]
BILLING CODE 4162–20–P

# 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 email 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—Existing Data Collection without an OMB Number—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID) (proposed), 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		Responses per respondent	Average burden per respondent (in hours)
School, school district or public health department	500	1	5/60

Dated: March 17, 2010.

#### Maryam I. Daneshvar,

Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. 2010-6523 Filed 3-23-10; 8:45 am]

BILLING CODE 4163-18-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

## Centers for Disease Control and Prevention

[30Day-10-0600]

## 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 email 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**

Model Performance Evaluation Program for Mycobacterium tuberculosis and Non-tuberculous Mycobacterium Drug Susceptibility Testing (OMB Control No. 0920–0600, expiration date 03/31/2010)—Revision—National Center for Preparedness, Detection, and Control of Infectious Diseases (NCPDCID), Centers for Disease Control and Prevention (CDC). Background and Brief Description

As part of the continuing effort to support both domestic and global public health objectives for treatment of tuberculosis (TB), prevention of multidrug resistance, and surveillance programs, CDC is requesting approval from the Office of Management and Budget to revise a currently approved data collection, the Model Performance Evaluation Program for *Mycobacterium* tuberculosis and Non-tuberculous Mycobacterium Drug Susceptibility Testing. This request includes changes to the Results Form and re-introduction of the Laboratory Practices Questionnaire.

While the overall number of cases of TB in the U.S. has decreased, rates still remain high among foreign-born persons, prisoners, homeless populations, and individuals infected with HIV in major metropolitan areas. The rate of TB cases detected in foreignborn persons has been reported to be more than nine times higher than the rate among the U.S. born population. CDC's goal to eliminate TB will be virtually impossible without considerable effort in assisting heavy disease burden countries in the reduction of tuberculosis. The Model Performance Evaluation Program for Mycobacterium tuberculosis and Nontuberculous Mycobacterium Drug Susceptibility Testing program supports this role by monitoring and evaluating the level of performance and practices among national and international laboratories performing M. tuberculosis susceptibility testing. Participation in this program is one way laboratories can ensure high-quality laboratory testing, resulting in accurate and reliable testing results.

By providing an evaluation program to assess the ability of the laboratories to test for drug resistant *M. tuberculosis* and selected strains of Non-tuberculous *Mycobacteria* (NTM), laboratories also have a self-assessment tool to aid in optimizing their skills in susceptibility testing. The information obtained from laboratories on susceptibility testing practices and procedures is used to establish variables related to good performance, assessing training needs, and aid with the development of practice standards.

Participants in this program include clinical and public health laboratories. Participants register by submitting an Enrollment Form. Data collection from domestic laboratory participants occurs twice per year. The data collected in this program will include the susceptibility test results of primary and secondary drugs, drug concentrations, and test methods performed by laboratories on a set of performance evaluation (PE) samples. The PE samples are sent to participants twice a year. Participants also report demographic data such as laboratory type and the number of tests performed annually. Participants report this data every two years. The burden for the Laboratory Practices Questionnaire has been adjusted for the average per year, since responses are received every other year. Participants may submit changes about their laboratory using the Laboratory Information Change Form.

There is no cost to respondents to participate other than their time. The total annualized burden for this information collection request is 166 hours.

### ESTIMATE OF ANNUALIZED BURDEN HOURS

Form	Respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Enrollment form	Labs	4 4 132 66	1 1 2 1	5/60 5/60 30/60 30/60