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.

There is no cost to respondents to participate other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Domestic Laboratory	Participant Biosafety Compliance Letter of Agreement.	93	2	5/60	16
	MPEP *Mycobacterium tuberculosis* Results Worksheet.	93	2	30/60	93
	Online Survey Instrument	93	2	15/60	47
Total			0		156

Dated: January 14, 2013.

Ron A. Otten,

Director, Office of Scientific Integrity (OSI), Office of the Associate Director for Science (OADS), Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2013–00988 Filed 1–17–13; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–13–0488]

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 (404) 639–7570 or send an email to *omb@cdc.gov.* Send written comments to 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

Restriction on Interstate Travel of Persons (OMB Control No. 0920–0488, Exp. 3/31/2013)—Revision—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention is requesting OMB approval for a revision of the information collection, ''Restriction on Interstate Travel of Persons'' (OMB Control No. 0920–0488).

This information collection request is scheduled to expire on March 31, 2013. CDC is authorized to collect this information under 42 CFR 70.5 (Certain communicable diseases; special requirements). This regulation requires that any person who is in the communicable period for cholera, plague, smallpox, typhus, or yellow fever or having been exposed to any such disease is in the incubation period thereof, to apply for and receive a permit from the Surgeon General or his authorized representative in order to travel from one State or possession to another.

CDC is requesting changes to the forms used within this information collection. The changes involve splitting the current form into two separate forms based on the type of respondent: an ill traveler, or the master of a vessel or conveyance engaged in interstate travel. CDC is also adding the option of electronic reporting of illness.

Control of disease transmission within the States is considered to be the province of state and local health authorities, with Federal assistance being sought by those authorities on a cooperative basis without application of Federal regulations. The regulations in 42 Part 70 were developed to facilitate Federal action in the event of large outbreaks requiring a coordinated effort involving several states, or in the event of inadequate local control. While it is not known whether, or to what extent situations may arise in which these regulations would be invoked, contingency planning for domestic emergency preparedness is now commonplace. Should these situations arise, CDC will use the reporting and recordkeeping requirements contained in the regulations to carry out quarantine responsibilities as required by law. The total number of burden hours requested for this collection is 3,701.

There is no cost to respondents other than their time.

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Traveler	42 CFR 70.3 Application to the State of destination for a permit.	2,000	1	15/60
Attending physician	42 CFR 70.3 Copy of material submitted by applicant and permit issued by State health authority.	2,000	1	15/60
State health authority	42 CFR 70.3 Copy of material submitted by applicant and permit issued by State health authority.	8	250	6/60

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Master of a vessel or person in charge of a conveyance.	42 CFR 70.4 Report by the master of a vessel or person in charge of conveyance of the incidence of a communicable disease occurring while in interstate travel (Paper Form if requested by CDC during public health emergency).	1,500	1	15/60
State health authority	42 CFR 70.4 Copy of material submitted to state or local health authority under this provision (Paper Form if requested by CDC during public health emergency).	20	75	6/60
Master of a vessel or person in charge of a conveyance.	42 CFR 70.4 Report by the master of a vessel or person in charge of conveyance of the incidence of a communicable disease occurring while in interstate travel (Radio or other telecommunication for routine reporting).	200	1	15/60
State health authority	42 CFR 70.4 Copy of material submitted to state or local health authority under this provision (Radio or other telecommunication for routine reporting).	200	1	15/60
Traveler	42 CFR 70.5 Application for a permit to move from State to State while in the communicable period.	3,750	1	15/60
Attending physician	42 CFR 70.5 Application for a permit to move from State to State while in the communicable period.	3,750	1	15/60

Dated: January 10, 2013.

Ron A. Otten,

Director, Office of Scientific Integrity (OSI), Office of the Associate Director for Science (OADS), Office of the Director Centers for Disease Control and Prevention.

[FR Doc. 2013–00987 Filed 1–17–13; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Office for State, Tribal, Local and Territorial Support (OSTLTS) Meeting; Correction

SUMMARY: This document corrects a notice that was published in the **Federal Register** on January 7, 2013 (78 FR 949–950). The 10th Biannual Tribal Consultation session has been postponed to coincide with the summer 2013 meetings; the dates will be announced once they are determined. The Tribal Advisory Committee Meeting will be extended and held February 5, 6, and 7, 2013, from 8:00 a.m.–4:30 p.m.

FOR FURTHER INFORMATION CONTACT: Kimberly Cantrell, Deputy Associate Director for Tribal Support, OSTLTS, via mail to 4770 Buford Highway NE., MS E–70, Atlanta, Georgia 30341 or email to *klw6@cdc.gov.*

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: January 11, 2013.

John Kastenbauer, J.D.,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2013–00989 Filed 1–17–13; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[CDC–2013–0001; NIOSH–134–B]

Update of NIOSH Nanotechnology Strategic Plan for Research and Guidance

AGENCY: National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Request for Information: Update of NIOSH Nanotechnology Strategic Plan for Research and Guidance.

SUMMARY: The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC) seeks comment on the types of hazard identification and risk management research that should be considered for updating the NIOSH FY2013–FY2016 nanotechnology strategic plan. This draft strategic plan (*Protecting the Nanotechnology Workforce: NIOSH Nanotechnology Research and Guidance Strategic Plan 2013–2016*) can be found in Docket CDC–2013–0001 at *http://www.regulations.gov.*

DATES: Comments must be received March 19, 2013.

ADDRESSES: You may submit comments, identified by CDC–2013–0001 and Docket Number NIOSH–134–B, by either of the two following methods:

- *Federal erulemaking portal: http://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* NIOSH Docket Office, Robert A. Taft Laboratories, MS–C34, 4676 Columbia Parkway, Cincinnati, OH 45226.

Instructions: All information received in response to this notice must include the agency name and docket number (CDC–2013–0001; NIOSH–134–B). All relevant comments received will be posted without change to *http://www.regulations.gov,* including any personal information provided. For access to prior background documents or previous comments received, go to *http://www.cdc.gov/niosh/docket/archive/docket134.html* and *http://www.cdc.gov/niosh/docket/archive/docket134A.html.*

FOR FURTHER INFORMATION CONTACT: Charles L. Geraci, NIOSH, Robert A. Taft Laboratories, MS–C14, 4676 Columbia Parkway, Cincinnati, Ohio 45226, telephone (513) 533–8339.

SUPPLEMENTARY INFORMATION:

Background

Since 2004, the National Institute for Occupational Safety and Health