these activities elicit the desired effects or impact.

ATSDR is requesting a reinstatement with minor changes for the following customer satisfaction surveys:

- ATSDR Web Site User Satisfaction Survey (WSUS)
- Toxicological Profiles User Satisfaction Survey (TPUS)
- ToxFAQs™ User Satisfaction Survey (TFUS)
- Public Health Statements User Satisfaction Survey (PHSUS)
- Toxicology Curriculum for Communities Training Manual User Satisfaction Survey (TCCUS)
- ToxProfiles™ CD–ROM User Satisfaction Survey (TP–CDUS)

The results of this project will ensure that these audiences will continue to find our knowledge products and informational pieces easy to access, clear, informative and useful. Specifically, this project will continue to examine whether current and future informational updates are presented in an appropriate technological format and whether they meet the needs, wants, and preferences of visitors to the ATSDR Web site. The survey questions have been held to the absolute minimum required for the use of the data.

This extension request is for a three-year period. There are no costs to the respondents other than their time. The total estimated annualized burden hours are 166.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
ATSDR Web site Visitors	WSUS	1000	1	5/60
	TPUS	300	1	5/60
	TFUS	300	1	5/60
	PHSUS	100	1	5/60
	TCCUS	160	1	5/60
	TP–CDUS	140	1	5/60

Dated: July 9, 2007.

Maryam I. Daneshvar,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. E7–13597 Filed 7–12–07; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–07–0274]

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) 395–6974. Written comments should be received within 30 days of this notice.

Proposed Project

CDC Model Performance Evaluation Program (MPEP) (0920–0274)—Revision—National Center for Preparedness, Detection, and Control of Infectious Diseases (proposed) (NCPDCID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

CDC is requesting OMB approval of a revision to its data collection, the CDC Model Performance Evaluation Program (MPEP). CDC originally implemented MPEP in 1986 to evaluate the performance of laboratories conducting testing to detect human immunodeficiency virus type 1 (HIV–1) antibody (Ab). CDC is requesting a 3-year approval for this data collection.

In this program, respondents receive 2 shipments of specimens per year. Respondents test the specimens in their laboratory/testing site and report their results either using a report booklet or on-line. CDC provides the respondent with a report containing the analysis of the laboratory test results reported to CDC. Participation in this program is voluntary and provides the respondents an opportunity to (1) assure accurate tests are being provided by the laboratory/testing site through external quality assessment; (2) improve testing quality through self-evaluation in a non-regulatory environment; (3) test well characterized samples from a source outside the test kit manufacturer; (4) discover potential testing problems so that procedures can be adjusted to eliminate them; (5) compare of testing results with others at a national and international level; and (6) consult with CDC staff to discuss testing issues.

In this request, CDC proposes to make the following revisions to the currently approved data collection:

- Addition of a Name and Address change form to report changes for the MPEP manager and coordinator at the respondent laboratory;
- Inclusion of additional test kit manufacturers approved by the FDA since previous OMB approval; and
- Elimination of reporting HIV–1 RNA Viral Load and CD4+ T-cell determinations.

There are no costs to the respondents other than their time to complete the survey. The total estimated annualized burden hours are 257.

TOTAL ESTIMATED ANNUALIZED BURDEN HOURS

Respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
New Enrollees	100	1	3/60
Laboratory Change Form	20	1	3/60

TOTAL ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Laboratory Test Result Form	754	2	10/60

Dated: July 9, 2007.

Maryam Daneshvar,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. E7–13605 Filed 7–12–07; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Committee on Childhood Lead Poisoning Prevention (ACCLPP), National Center for Environmental Health (NCEH)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announce the following meeting of the aforementioned committee:

Times and Dates:
8:30 a.m.–5 p.m., September 18, 2007.
8:30 a.m.–12:30 p.m., September 19, 2007.

Place: Radisson Plaza Hotel Minneapolis, 35 South 7th Street, Minneapolis, MN 55402, Telephone: (612) 339–4900.

Open to the public, limited only by the space available. The meeting room accommodates approximately 75 people.

Purpose: The Committee provides advice and guidance to the Secretary; the Assistant Secretary for Health; and the Director, CDC, regarding new scientific knowledge and technological developments and their practical implications for childhood lead poisoning prevention efforts. The Committee also reviews and reports regularly on childhood lead poisoning prevention practices and recommends improvements in national childhood lead poisoning prevention efforts.

Matters To Be Discussed: The meeting will include discussion on the potential approaches to strengthen existing strategies to achieve the Healthy People 2010 goal of eliminating Elevated Blood Lead Levels as a Public Health Problem in the United States by 2010, the development of a prevention based research agenda, and the study designs related to adverse effects from Blood Lead Levels (BLLs) < 10 µg/dl; and updates on the school performance and concurrent BLLs.

Agenda items are subject to change as priorities dictate.

Opportunities will be provided during the meeting for oral comments. Depending on the time available and the number of requests, it may be necessary to limit the time of each presenter.

Contact Person for More Information: Claudine Johnson, Clerk, Lead Poisoning Prevention Branch, Division of Environmental Emergency Health Services, NCEH, CDC, 4770 Buford Hwy, NE., Mailstop F–40, Atlanta, GA 30341, Telephone (770) 488–3629, Fax (770) 488–3635.

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 CDC and the Agency for Toxic Substances and Disease Registry.

Dated: July 6, 2007.

Elaine L. Baker,

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

[FR Doc. E7–13596 Filed 7–12–07; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel: Racial and Ethnic Approaches to Community Health Across the United States (REACH US), Request for Applications (RFA) DP07–707

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Public Law 92–463), the Centers for Disease Control and Prevention (CDC) announces a meeting of the aforementioned Special Emphasis Panel.

Times and Dates: 3 p.m.–8 p.m., July 31, 2007 (Closed), 8 a.m.–5 p.m., August 1, 2007 (Closed), 8 a.m.–5 p.m., August 2, 2007 (Closed), 3 p.m.–8 p.m., August 6, 2007 (Closed), 8 a.m.–5 p.m., August 7, 2007 (Closed), 8 a.m.–5 p.m., August 8, 2007 (Closed), 8 a.m.–5 p.m., August 9, 2007 (Closed).

Place: Sheraton Gateway Hotel Atlanta Airport, 1900 Sullivan Road, Atlanta, GA 30337, Telephone (800) 454–6835.

Status: The meeting will be closed to the public in accordance with provisions set forth in section 552b(c)(4) and (6), Title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Pub. L. 92–463.

Matters To Be Discussed: The meeting will include the review, discussion, and evaluation of research grant applications in response to RFA DP07–707, ''Racial and Ethnic Approaches to Community Health Across the United States (REACH US).''

For Further Information Contact: Thijuanie Lockhart, Program & Management Analyst, CDC, 4770 Buford Highway, NE., Mail Stop K–30, Atlanta, GA 30341, Telephone (404) 488–5303.

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 CDC and the Agency for Toxic Substances and Disease Registry.

Dated: July 9, 2007.

Elaine L. Baker,

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

[FR Doc. E7–13602 Filed 7–12–07; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Statement of Organization, Functions, and Delegations of Authority

Part C (Centers for Disease Control and Prevention) of the Statement of Organization, Functions, and Delegations of Authority of the Department of Health and Human Services (45 FR 67772–67776, dated October 14, 1980, and corrected at 45 FR 69296, October 20, 1980, as amended most recently at 72 FR 35491–35492, dated June 28, 2007) is amended to reflect the establishment of the Extramural Research Program Office within the National Center for Chronic Disease Prevention and Health Promotion, Coordinating Center for Health Promotion, Centers for Disease Control and Prevention.

Section C–B, Organization and Functions, is hereby amended as follows: After the functional statement for the Program Services Branch (CUC13), Office of the Director (CUC1), National Center for Chronic Disease Prevention and Health Promotion (CUC), insert the following:

Extramural Research Program Office (CUC18). The Extramural Research Program Office (ERPO) plans, develops,