

Questionnaires; (3) buccal cell collection; (4) a telephone interview of the biological mother and/or primary caregiver; (5) a child development evaluation (more comprehensive for case participants than for the control

group participants); (6) parent-child development interview (for case participants only); (7) a physical exam of the child; (8) biological sampling of the child (blood and hair); and (9)

biological sampling of the biological parents (blood only).

There are no costs to respondents other than their time. The total estimated annualized burden is 30,103 hours.

#### ESTIMATED ANNUALIZED BURDEN HOURS

Respondents	Number of respondents	Number of responses per respondent	Average burden per response (In hrs)	Total burden hours
Contact by Mail .....	17,610	1	10/60	2,935
Telephone Contact .....	8,922	1	20/60	2,974
Parent Questionnaires and biologic sample .....	3,456	1	235/60 (3h 55m)	13,548
Caregiver Interview .....	3,282	1	30/60	1,641
Clinic Visit (Child Development Evaluation, physical exam, and biosamples):				
• Case .....	810	1	355/60 (5h 55m)	4,793
• NIC .....	1,170	1	110/60 (1h 50m)	2,145
• Subcohort .....	1,134	1	110/60 (1h 50m)	2,079
TOTAL .....				30,103

Dated: December 26, 2006.

**Joan F. Karr,**

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

[FR Doc. E6-22579 Filed 1-4-07; 8:45 am]

**BILLING CODE 4163-18-P**

#### DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### Centers for Disease Control and Prevention

[30Day-07-05CG]

#### 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](mailto: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

Morbidity Monitoring Project (MMP)—New—National Center for HIV, STD and TB Prevention (NCHSTP), Centers for Disease Control and Prevention (CDC).

#### Background and Brief Description

The Morbidity Monitoring Project (MMP) is a new project. The purpose of MMP is to supplement the HIV/AIDS surveillance programs in 26 selected state and local health departments, which collect information on persons diagnosed with, living with, and dying from HIV infection and AIDS and will incorporate data elements from two data collections: Supplement to HIV/AIDS Surveillance (SHAS) project (0920-0262) and the Adult/Adolescent Spectrum of HIV Disease (ASD).

MMP will collect data on behaviors and clinical outcomes from a probability sample of HIV-infected adults receiving care in the U.S. Collection of data from interviews with HIV-infected patients will provide information on patient demographics, and the current levels of behaviors that may facilitate HIV transmission: sexual and drug use behaviors; patients' access to, use of and barriers to HIV-related secondary

prevention services; utilization of HIV-related medical services; and adherence to drug regimens. Collection of data from patient medical records will provide information on: demographics and insurance status; the prevalence and incidence of AIDS-defining opportunistic illnesses and comorbidities related to HIV disease; the receipt of prophylactic and antiretroviral medications; and whether patients are receiving screening and treatment according to Public Health Service guidelines. No other Federal agency collects national population-based behavioral and clinical information from HIV-infected adults in care.

The data will have significant implications for policy, program development, and resource allocation at the state/local and national levels. Users of MMP data include, but are not limited to, Federal agencies, state and local health departments, clinicians, researchers, and HIV prevention and care planning groups. There are no costs to the respondents other than their time. The total estimated annualized burden hours are 6,101.

#### ESTIMATED ANNUALIZED BURDEN HOURS

Respondents	Number of respondents	Number of responses per respondent	Average burden per response (In hours)
Persons interviewed with standard interview .....	7,988	1	45/60
Persons interviewed with short interview .....	166	1	20/60

## ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Respondents	Number of respondents	Number of responses per respondent	Average burden per response (In hours)
Persons interviewed with proxy interview .....	166	1	20/60

Dated: December 26, 2006.

**Joan F. Karr,**

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

[FR Doc. E6–22600 Filed 1–4–07; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

#### National Center for Environmental Health/Agency for Toxic Substances and Disease Registry

The Health Department Subcommittee of the Board of Scientific Counselors (BSC), Centers for Disease Control and Prevention (CDC), National Center for Environmental Health (NCEH)/Agency for Toxic Substances and Disease Registry (ATSDR): Teleconference Meeting.

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention, NCEH/ATSDR announces the following teleconference meeting of the aforementioned subcommittee:

*Times and Dates:* 12:30 p.m.–2 p.m., February 12, 2007.

*Place:* Century Center, 1825 Century Boulevard, Atlanta, Georgia 30345.

*Status:* Open to the public, teleconference access limited only by availability of telephone ports.

*Purpose:* Under the charge of the BSC, NCEH/ATSDR, the Health Department Subcommittee will provide the Board with advice and recommendations on local and state health department issues and concerns that pertain to the mandates and mission of NCEH/ATSDR.

*Matters to be Discussed:* The meeting agenda will include a review of agenda items and approval of minutes; bridging NCEH/ATSDR programs; public comment; and the next steps for the Health Department Subcommittee.

Items are subject to change as priorities dictate.

*Supplementary Information:* This teleconference meeting is scheduled to begin at 12:30 p.m. Eastern Standard Time. To participate, dial (877) 315–6535 and enter conference code 383520. The public comment period will be held from 1:50 p.m.–2 p.m.

*For Further Information Contact:* Please contact Shirley D. Little, Committee Management Specialist, NCEH/ATSDR, 1600 Clifton Road, Mail Stop E–28, Atlanta, GA 30303; telephone 404/498–0003, fax 404/498–0059; e-mail: [slittle@cdc.gov](mailto:slittle@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 CDC and the National Center for Environmental Health/Agency for Toxic Substances and Disease Registry.

Dated: December 28, 2006.

**Elaine Baker,**

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

[FR Doc. E6–22585 Filed 1–4–07; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

[Document Identifier: CMS–R–193]

#### Agency Information Collection Activities: Proposed Collection; Comment Request

**AGENCY:** Centers for Medicare & Medicaid Services, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare & Medicaid Services (CMS) is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (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:* Revision of a currently

approved collection; *Title of Information Collection:* Important Message from Medicare Use: Requirements that hospitals notify beneficiaries in inpatient hospital settings of their rights as a hospital patient including their discharge appeal rights are referenced in Section 1866(a)(1)(M) of the Social Security Act. The authority for the right to an expedited determination is set forth at Section 1869(c)(3)(C)(iii)(III) of the Act. Under sections 42 CFR 405.1205 and 422.620, the hospital must deliver valid, written notice, the Important Message from Medicare (IM), of a patient's rights as a hospital patient including the discharge appeal rights, within 2 calendar days of admission. A follow-up copy of the signed IM is given again as far as possible in advance of discharge, but no more than 2 calendar days before. Follow-up notice is not required if the provision of the admission IM, falls within 2 calendar days of discharge. *Form Number:* CMS–R–193 (OMB #: 0938–0692); *Frequency:* Reporting: Yearly; *Affected Public:* Business or other for-profit and Not-for-profit institutions; *Number of Respondents:* 6,000; *Total Annual Responses:* 13,000,000; *Total Annual Hours:* 2,990,000.

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](mailto: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 at the address below, no later than 5 p.m. on March 6, 2007.

CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development—C, Attention: Bonnie L Harkless, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.