

Estimated Annual Costs to the Federal Government

The total cost to the government for activities directly related to this collection is \$432,451,000.

Request for Comments

In accordance with the above cited legislation, comments on the AHRQ information collection proposal are requested with regard to any of the following: (a) Whether the proposed collection of information is necessary for the proper performance of functions of the AHRQ, including whether the information will have practical utility; (b) the accuracy of the AHRQ's estimate of the burden (including hours and costs) 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 the use of automated collection techniques or other forms of information technology.

Comments submitted in response to this notice will be summarized and included in the request for OMB approval of the proposed information collection. All comments will become a matter of public record.

Dated: February 27, 2003.

Carolyn M. Clancy,

Director

[FR Doc. 03-5298 Filed 3-5-03; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Committee on Childhood Lead Poisoning Prevention (ACCLPP): Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the National Center for Environmental Health (NCEH) of the Centers for Disease Control and Prevention (CDC) announces the following committee meeting.

Name: Advisory Committee on Childhood Lead Poisoning Prevention.

Time and Date: 8:30 a.m.–5:30 p.m., March 18, 2003.

Place: Hilton—Crystal City at National Airport, 2399 Jefferson Davis Highway, Arlington, VA 22202, telephone 703/418-6800.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 55 people.

Purpose: The Committee shall provide 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 shall also review and report regularly on childhood lead poisoning prevention practices and recommend improvements in national childhood lead poisoning prevention efforts.

Matters to be Discussed: Agenda items include: Updates on Primary Prevention issues, Medicaid Targeted Screening, Review of Evidence for Effects at Blood Lead Levels <10 µg/dL issues, Screening of Immigrant/Adopted Children, and Study of Relationship of Environmental Tobacco Smoke and Blood Lead Levels.

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: Crystal M. Gresham, Program Analyst, Lead Poisoning Prevention Branch, Division of Emergency and Environmental Health Services, NCEH, CDC, 1600 Clifton Road, NE., M/S F-30, Atlanta, Georgia 30333, telephone 770/488-7490, fax 770/488-4178.

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: February 28, 2003.

Alvin Hall,

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

[FR Doc. 03-5247 Filed 3-5-03; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Revised Vaccine Information Materials for Measles, Mumps and Rubella Vaccines; Revised Instructions for Use of Vaccine Information Statements

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: Under the National Childhood Vaccine Injury Act (42 U.S.C. 300aa-26), the CDC must develop vaccine information materials that all health care providers are required to give to patients/parents prior to administration of specific vaccines. Since the recommended interval

between receiving rubella-containing vaccine and becoming pregnant has been amended from 3 months to 4 weeks, the vaccine information materials covering measles, mumps and rubella vaccine needed to be revised. On October 10, 2002, CDC published a notice in the **Federal Register** (67 FR 63106) seeking public comments on the proposed revised vaccine information materials for measles, mumps and rubella vaccines. The 60 day comment period ended on December 9, 2002. Following review of the comments submitted and consultation as required under the law, CDC has finalized these vaccine information materials. The final materials, and revised instructions for their use and for use of materials for other covered vaccines, are contained in this notice.

DATES: Beginning as soon as practicable, each health care provider who administers any vaccine that contains measles, mumps or rubella vaccine shall, prior to administration of each dose of the vaccine, provide a copy of the vaccine information materials contained in this notice, dated January 15, 2003, to the parent or legal representative of any child to whom such provider intends to administer the vaccine and to any adult to whom such provider intends to administer the vaccine, in lieu of providing earlier versions of these materials.

FOR FURTHER INFORMATION CONTACT:

Walter A. Orenstein, M.D., Director, National Immunization Program, Centers for Disease Control and Prevention, Mailstop E-05, 1600 Clifton Road, NE., Atlanta, Georgia 30333, telephone (404) 639-8200.

SUPPLEMENTARY INFORMATION: The National Childhood Vaccine Injury Act of 1986 (Pub. L. 99-660), as amended by section 708 of Public Law 103-183, added section 2126 to the Public Health Service Act. Section 2126, codified at 42 U.S.C. 300aa-26, requires the Secretary of Health and Human Services to develop and disseminate vaccine information materials for distribution by all health care providers in the United States to any patient (or to the parent or legal representative in the case of a child) receiving vaccines covered under the National Vaccine Injury Compensation Program.

Development and revision of the vaccine information materials have been delegated by the Secretary to the Centers for Disease Control and Prevention (CDC). Section 2126 requires that the materials be developed, or revised, after notice to the public, with a 60-day comment period, and in consultation with the Advisory Commission on