DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Presidential Advisory Council on HIV/AIDS

AGENCY: Department of Health and Human Services, Office of the Secretary, Office of Public Health and Science.

ACTION: Notice.

SUMMARY: As stipulated by the Federal Advisory Committee Act, the U.S. Department of Health and Human Services (DHHS) is hereby giving notice that the Presidential Advisory Council on HIV/AIDS (PACHA) will hold a meeting. The meeting will be open to the public.

DATES: The meeting will be held on Monday, April 26, 2010 from 10 a.m. to approximately 4 p.m. The meeting will be open to the public.

ADDRESSES: The White House, South West Auditorium, Eisenhower Executive Office Building, State Avenue and 17th Street, NW., Washington, DC.

FOR FURTHER INFORMATION CONTACT: Mr. Melvin Joppy, Committee Manager, Presidential Advisory Council on HIV/AIDS, Department of Health and Human Services, 200 Independence Avenue, SW., Room 443H, Hubert H. Humphrey Building, Washington, DC 20210; (202) 690–5560. More detailed information about PACHA can be obtained by accessing the Council’s Web site at http://www.pacha.gov.

SUPPLEMENTARY INFORMATION: PACHA was established by Executive Order 12963, dated June 14, 1995 as amended by Executive Order 13009, dated June 14, 1996. The Council was established to provide advice, information, and recommendations to the Secretary regarding programs and policies intended to (a) Promote effective prevention of HIV disease, (b) advance research on HIV and AIDS, and (c) promote quality services to persons living with HIV disease and AIDS. PACHA was established to serve solely as an advisory body to the Secretary of Health and Human Services.

The agenda for this Council meeting will be posted on the Council’s website http://www.pacha.gov.

This meeting of the PACHA will be on White House property, thus, each person must be screened and cleared by the U.S. Secret Service. Pre-registration for public attendance is mandatory. Please contact: Natalie Pojman, Office of National AIDS Policy (202) 456–4533 or npojman@who.eop.gov. Members of the public will be accommodated on a first come first served basis as meeting room space is limited. Ms. Pojman will need your full name, social security number, date of birth, residency, and country of origin to process public access attendance. Pre-registration must be submitted by close of business Thursday, April 22, 2010.

Members of the public will have the opportunity to provide comments at the meeting. If you plan to make a public comment you must pre-register with Natalie Pojman, Office of National AIDS Policy. Public comments will be limited to two minutes per speaker. Any members of the public who wish to have printed material distributed to PACHA members for discussion at the meeting should submit, at a minimum, 30 copies of the materials to the Committee Manager, PACHA, no later than close of business April 22, 2010. Contact information for the PACHA Committee Manager is listed above. Justification for filing notice less than 15 days prior to meeting: PACHA meetings are scheduled to be held in coordination with the White House Office of National AIDS Policy (ONAP), which partners with the HHS Office of HIV/AIDS Policy to provide management oversight for the Council’s activities. Meeting dates are selected in consideration of the availability of meeting space and ONAP staff attendance. The designated date was recently identified because both the desired meeting site and ONAP staff are available.

Dated: April 8, 2010.

Christopher Bates, Executive Director, Presidential Advisory Council on HIV/AIDS.

[FR Doc. 2010–8548 Filed 4–13–10; 8:45 am]

BILLING CODE 4150–43–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–10–0733]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and
Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404–639–5960 and send comments to Maryam I. Daneshvar, CDC Acting Reports Clearance Officer, 1600 Clifton Road, MS–D74, Atlanta, GA 30333 or send an e-mail to omb@cdc.gov.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency’s estimate of the burden 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 through the use of automated collection techniques or other forms of information technology. Written comments should be received within 60 days of this notice.

**Proposed Project**


**Background and Brief Description**

The National Center on Birth Defects and Developmental Disabilities at CDC promotes the health of babies, children, and adults with disabilities. As part of these efforts the Center is actively involved in addressing hearing loss (HL) among newborns and infants. HL is a common birth defect that affects approximately 12,000 infants each year and, when left undetected, can result in developmental delays. As awareness about infant HL increases, so does the demand for accurate information about rates of screening, referral, loss to follow-up, and incidence. This information is important for helping to ensure infants and children are receiving recommended screening and follow-up services, documenting the occurrence and etiology of differing degrees of HL among infants, and determining the overall impact of infant HL on future outcomes, such as cognitive development, and family dynamics. These data will also assist state Early Hearing Detection and Intervention (EHDI) programs with quality improvement activities and provide information that will be helpful in assessing the impact of federal initiatives. The public will be able to access this information via the CDC EHDI Web site (http://www.cdc.gov/ncbddd/ehdi/data.htm).

Given the lack of a standardized and readily accessible source of data, the CDC EHDI program developed a survey to be used annually that utilizes uniform definitions to collect aggregate, standardized EHDI data from states and territories. The request to complete this survey is planned to be disseminated to respondents via an e-mail, which will include a summary of the request and other relevant information. Minor changes to this survey, based on respondent feedback, are planned in order to make the survey easier to complete and further improve data quality. These changes include splitting the previously combined questions about the number of infants that died and parents refused into two separate questions, adding a question about how many infants with hearing loss are receiving only monitoring services, simplifying the table for reporting type and severity of hearing loss data, and expanding the maternal race categories in the demographic section.

There are no costs to the respondents other than their time.

**Estimated Annualized Burden Hours**

<table>
<thead>
<tr>
<th>Respondents</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden per response (in hours)</th>
<th>Total burden (in hours)</th>
</tr>
</thead>
<tbody>
<tr>
<td>State and territory EHDI Program Coordinators: Those who review survey instructions ..................................................</td>
<td>57</td>
<td>1</td>
<td>10/60</td>
<td>10</td>
</tr>
<tr>
<td>State and territory EHDI Program Coordinators: Those who complete the survey ..................................................</td>
<td>50</td>
<td>1</td>
<td>4</td>
<td>200</td>
</tr>
</tbody>
</table>

**SUMMARY:** The Administration on Aging (AoA) is announcing that the proposed collection of information listed below has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

**DATES:** Submit written comments on the collection of information by May 14, 2010.

**ADDRESSES:** Submit written comments on the collection of information by fax 202.395.6974 to the OMB Desk Officer for AoA, Office of Information and Regulatory Affairs, OMB.

**FOR FURTHER INFORMATION CONTACT:** Valerie Cook at 202–357–3583

**SUPPLEMENTARY INFORMATION:** In compliance with 44 U.S.C. 3507, AoA has submitted the following proposed collection of information to OMB for review and clearance.

The Older Americans Act (OAA) requires annual program performance reports from States. In compliance with this OAA provision, AoA developed a State Program Report (SPR) in 1996 as part of its National Aging Program Information System (NAPIS). The SPR collects information about how State Agencies on Aging expend their OAA funds as well as funding from other sources for OAA authorized supportive services. The SPR also collects information on the demographic and functional status of the recipients, and is a key source for AoA performance measurement. This collection includes minor revisions of the format from the 2006 approved version. The proposed revised version will be in effect for the