classes of workers to the Special Exposure Cohort (SEC).

In December 2000, the President delegated responsibility for funding, staffing, and operating the Advisory Board to HHS, which subsequently delegated this authority to CDC. NIOSH implements this responsibility for CDC. The charter was issued on August 3, 2001, renewed at appropriate intervals, and will expire on August 3, 2011.

Purpose: This Advisory Board is charged with (a) providing advice to the Secretary, HHS, on the development of guidelines under Executive Order 13179; (b) providing advice to the Secretary, HHS, on the scientific validity and quality of dose reconstruction efforts performed for this program; and (c) upon request by the Secretary, HHS, advising the Secretary on whether there is a class of employees at any Department of Energy (DOE) facility who were exposed to radiation but for whom it is not feasible to estimate their radiation dose, and on whether there is reasonable likelihood that such radiation doses may have endangered the health of members of this class.

Matters To Be Discussed: The agenda for the Advisory Board meeting includes: NIOSH Program Update; Department of Labor (DOL) Program Update; DOE Program Update; NIOSH Program Evaluation; HHS Proposed Rule to Amend Probability of Causation Guidelines Regarding Chronic Lymphocytic Leukemia (42 CFR pt. 81); Savannaĥ River Site Work Group Update; Feed Materials Production Center Work Group Update; Weldon Spring Work Group Update; SEC petitions for: Piqua Organic Moderated Reactor (1963–1966), Šandia National Laboratory (1957-1962), Hanford (Plutonium Finishing Plant, 1987–1989), General Electric (Evendale, Ohio); SEC Petition Status Updates; Subcommittee and Work Group Reports; Board Work Sessions, and an Administrative Session.

The agenda is subject to change as priorities dictate.

In the event an individual cannot attend, written comments may be submitted in accordance with the redaction policy provided below. Any written comments received will be provided at the meeting and should be submitted to the contact person below well in advance of the meeting.

Policy on Redaction of Board Meeting Transcripts (*Public Comment*): (1) If a person making a comment gives his or her name, no attempt will be made to redact that name. (2)

NIOSH will take reasonable steps to ensure that individuals making public comment are aware of the fact that their comments (including their name, if provided) will appear in a transcript of the meeting posted on a public Web site. Such reasonable steps include: (a) A statement read at the start of each public comment period stating that transcripts will be posted and names of speakers will not be redacted; (b) A printed copy of the statement mentioned in (a) above will be displayed on the table where individuals sign up to make public comments; (c) A statement such as outlined in (a) above will also appear with the agenda for a Board Meeting when it is posted on the NIOSH Web site; (d) A statement such as in (a) above will appear in the Federal Register Notice that announces Board and Subcommittee meetings. (3) If an individual in making a statement reveals personal information (e.g., medical information) about themselves that information will not usually be redacted. The NIOSH Freedom of Information Act (FOIA) coordinator will, however, review such revelations in accordance with the FOIA and the Federal Advisory Committee Act (FACA) and if deemed appropriate, will redact such information. (4) All disclosures of information concerning third parties will be redacted. (5) If it comes to the attention of the Designated Federal Officer that an individual wishes to share information with the Board but objects to doing so in a public forum, the DFO will work with that individual, in accordance with the FACA, to find a way that the Board can hear such comments.

Contact Person for More Information: Theodore Katz, M.P.A., Executive Secretary, NIOSH, CDC, 1600 Clifton Road, Mailstop E– 20, Atlanta Georgia 30333, telephone: (513) 533–6800, toll free: 1 (800)CDC–INFO, email: dcas@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.

**ANNUAL BURDEN ESTIMATES** 

Dated: April 29, 2011. Elaine L. Baker, Director, Management Analysis and Services Office, Centers for Disease Control and Prevention. [FR Doc. 2011–11076 Filed 5–5–11; 8:45 am] BILLING CODE 4163–18–P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

[OMB No. 0970-0278]

# Submission for OMB Review; Comment Request; Reunification Procedures for Unaccompanied Alien Children

#### Description

Following the passage of the 2002 Homeland Security Act (Pub. L 107– 296), the Administration for Children and Families (ACF), Office of Refugee Resettlement (ORR), is charged with the care and placement of unaccompanied alien children in Federal custody, and implementing a policy for the release of these children, when appropriate, upon the request of suitable sponsors while awaiting immigration proceedings. In order for ORR to make determinations regarding the release of these children, the potential sponsors must meet certain conditions pursuant to section 462 of the Homeland Security Act and the Flores v. Reno Settlement Agreement No. CV85 4544-RJK (C.D. Cal. 1997). The proposed information collection requests information to be utilized by ORR for determining the suitability of a sponsor/respondent for the release of a minor from ORR custody. The proposed instruments are the Sponsors Agreement to Conditions of Release, Verification of Release, Family Reunification Packet, and the Authorization for Release of Information.

*Respondents:* Sponsors requesting release of unaccompanied alien.

#### Number of Average Number of Total burden Instrument burden hours responses per respondents hours respondent per response Verification of Release (UAC) ..... 4,595 0.25 1148.75 1 Authorization for Release of Information (Sponsor) ..... 4,595 1 0.25 1,148.75 Family Reunification Packet (Sponsor) 4,595 1 4,595 1 Sponsors Agreement to Conditions of Release (Sponsor) ..... 4,595 0.25 1,148.75 1 Verification of Release (Case Worker) 4,595 0.25 1,148.75 1 Authorization for Release of Information (Case Worker) ..... 4,595 0.25 1 Family Reunification Packet (Case Worker) ..... 4.595 4,595 1 1 Sponsors Agreement to conditions of Release (Case Worker) ..... 0.25 4,595 1 1148.75

*Estimated Total Annual Burden Hours:* 16,082.50.

# **Additional Information**

Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. E-mail address: infocollection@acf.hhs.gov.

OMB Comment:

OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, *Fax:* 202–395–7285, *E-mail:* 

OIRA\_SUBMISSION@OMB.EOP.GOV. Attn: Desk Officer for the

Administration for Children and Families.

Dated: March 29, 2011. **Robert Sargis,** *Reports Clearance Officer.* 

[FR Doc. 2011–11046 Filed 5–5–11; 8:45 am] BILLING CODE 4184–01–P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

# Administration for Children and Families

### Proposed Information Collection Activity; Comment Request

*Title:* Continued Tracking of Families in the Head Start Impact Study. *OMB No.:* 0970–0229.

Description: The Administration for Children and Families (ACF) of the Department of Health and Human Services (HHS), is requesting comments on plans to collect information from children and families in the Head Start Impact Study. In anticipation of the possibility of conducting a follow-up for this study in early adulthood, this effort will collect information necessary to identify respondents' current location, as well as other basic information about the parents' whereabouts and future contacts, should the follow-up study be continued. A limited set of items will also be collected to gather information from parents about their children's wellbeing, including whether they have been retained in grade, are receiving special education services, how well they are faring in school, and how they behave. This information will be collected annually, with the goal of ensuring continued high response rates in future follow-up data collections.

The Head Start Impact Study was a longitudinal study that involved approximately 5,000 first time enrolled

# ANNUAL BURDEN ESTIMATES

Instrument	Annual number of respondents	Number of re- sponses per respondent	Average burden hours per response	Total annual burden hours
Tracking Interview	4,667	1	2/3	1,556

# *Estimated Total Annual Burden Hours:* 1,556.

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: OPRE Reports Clearance Officer. E-mail address: OPREinfocollection@acf.hhs.gov. All

requests should be identified by the title of the information collection.

The Department specifically requests comments 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) 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. Consideration will be given to

three- and four-year old preschool children across 84 nationally representative grantee/delegate agencies (in communities where there were more eligible children and families than can be served by the program.) The participating children were randomly assigned to either a Head Start group (that could enroll in Head Start services) or a control group (that could not enroll in Head Start services but could enroll in other available services selected by their parents). Data collection for the study began in fall of 2002 and extended through spring 2008, through the children's 3rd grade year. Tracking of these children and families has continued through spring 2011.

It is the intention of the Administration for Children and Families to continue to examine outcomes for this sample of children and families when the children reach early adulthood. In order to ensure that participants can be located for that future study, location and contact information will be collected from parents or guardians in the spring of 2012, 2013, 2014, 2015, and 2016. The tracking updates will primarily be conducted over the telephone with inperson follow-up as necessary. Tracking updates will take about 20 minutes to complete.

Respondents:

comments and suggestions submitted within 60 days of this publication.

Dated: May 2, 2011.

### Seth F. Chamberlain

Reports Clearance Officer. [FR Doc. 2011–11100 Filed 5–5–11; 8:45 am]

BILLING CODE 4184-22-P