program. 3001 children and families in 17 sites were randomly assigned either to the program group (allowed to enroll in EHS), or to the control group (precluded from enrolling in EHS, although they could receive other services in the community). Child and family assessments were conducted when children were 14 months old, 24 months old, 36 months old, in the spring prior to kindergarten entry, and again in the spring of the sixth year of formal schooling (5th grade for most children).

If the decision is made to follow the sample through high school, it is important to maintain contact with the participants so that response rates at follow-up points will be maximized and will not further deteriorate. The success that the contractor has in contacting the sample members will be taken into consideration in the decision whether to pursue further follow up. It is hoped that this strenuous effort in location and tracking will re-establish contact with participants who were not reached in the fifth grade data collection wave. Telephone interviews will be conducted in order to update the respondent's location and contact information. This information will be collected from parents or guardians in the spring of 2011. A small set of additional items will provide information on the parents' perception of the children's status.

Respondents: Treatment and control group members in the Early Head Start Research and Evaluation Project.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Tracking interview	2,700	1	0.25	675

Estimated Total Annual Burden Hours: 675.

In compliance with the requirements of Section 506(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 Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. E-mail address: *infocollection@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 comments and suggestions submitted within 60 days of this publication.

Dated: October 28, 2010.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2010–27593 Filed 11–1–10; 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

Proposed Projects:

Title: Reunification Procedures for Unaccompanied Alien Children.

OMB No.: 0970–0278.

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 children to their custody.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Sponsor's Agreement to Conditions of Release	4,288	2	0.08	686.08
Verification of Release	4,288	1	0.17	728.96
Family Reunification Packet	4,288	18	0.04	3,087.36
Authorization for Release of Information	4,288	15	0.22	14,150.40

Estimated Total Annual Burden Hours: 18,652.80.

In compliance with the requirements of Section 506(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 Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. E-mail address: *infocollection@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 comments and suggestions submitted within 60 days of this publication.

Dated: October 28, 2010.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2010–27596 Filed 11–1–10; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Diabetes and Digestive and Kidney Diseases; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel, Ancillary Study in Necrotizing Enterocolitis.

Date: December 1, 2010.

Time: 3 p.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892. (Telephone Conference Call)

Contact Person: Maria E. Davila-Bloom, PhD, Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 758, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 594–7637, *davila-bloomm@extra.niddk.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.847, Diabetes, Endocrinology and Metabolic Research; 93.848, Digestive Diseases and Nutrition Research; 93.849, Kidney Diseases, Urology and Hematology Research, National Institutes of Health, HHS)

Dated: October 26, 2010.

Jennifer S. Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010–27569 Filed 11–1–10; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No FDA–2010–N–0001]

Pediatric Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committee: Pediatric Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA's regulatory issues.

Date and Time: The meeting will be held on December 7, 2010, from 8 a.m. to 6 p.m.

Location: Bethesda Marriott Hotel, 5151 Pooks Hill Rd., Bethesda, MD 20814.

Contact Person: Walter Ellenberg, Office of Pediatric Therapeutics, Office of Special Medical Programs, Office of the Commissioner, Food and Drug Administration, 10903 New Hampshire Ave., Bldg 32, rm. 5154, Silver Spring, MD 20993, 301–796–0885, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 8732310001. Please call the Information Line for up-to-date information on this meeting. A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the agency's Web site and call the appropriate advisory committee hot line/phone line to learn about possible modifications before coming to the meeting.

Agenda: The Pediatric Advisory Committee will meet to discuss pediatric-focused safety reviews, as mandated by the Best Pharmaceuticals for Children Act and the Pediatric Research Equity Act, for Prevista (darunavir ethanolate), PegIntron (peginterferon alfa–2b), Xyzal (levocetirizine dihydrochloride) tablet and solution, Flovent HFA (fluticasone propionate), Acanya Gel (clindamycin/benzoyl peroxide combination), Epiduo Gel (adapalene and benzoyl peroxide), Ulesfia Lotion 5% (benzyl alcohol), Axert (almotriptan), Gardasil (human papillomavirus quadrivalent types 6, 11, 16, 18, vaccine recombinant), Lamictal and Lamictal XR (lamotrigine), and Neulasta (pegfilgrastim). The committee will also receive a followup on Depakote ER (divalproex sodium). Committee members who participated in the Cardiovascular and Renal Drugs Advisory Committee and the Gastrointestinal Drugs Advisory Committee meetings held on July 29, 2010, and November 5, 2010, respectively, will provide a brief summary of the meetings.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at *http://www.fda.gov/AdvisoryCommittees/Calendar/default.htm.* Scroll down to the appropriate advisory committee link.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before November 29, 2010. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. Those desiring to make formal oral presentations should notify the contact person and submit a brief