both the adult and pediatric groups. There are no costs to respondents. The total response burden for the study 201 hours.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents (adult and pediatric)	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Case Subjects > 17 years of age	Screening Process	129	1	5/60
	Telephone interview	71	1	30/60
Control Subjects > 17 years of age	Screening Process	142	1	5/60
	Telephone interview	71	1	30/60
Case Subject ≤ 1–5 years of age	Screening Process	141	1	5/60
	Telephone interview	78	1	30/60
Control Subjects ≤ 1–5 years of age	Screening Process	194	1	5/60
	Telephone interview	78	1	30/60
Total				

Leroy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2014–03013 Filed 2–11–14; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Head Start Family and Child Experiences Survey (FACES)

OMB No.: 0970–0151

Description: The Office of Planning, Research and Evaluation (OPRE), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), is proposing to collect data for a new round of the Head Start Family and Child Experiences Survey (FACES). Featuring a new ''Core Plus'' study design, FACES will provide data on a set of key indicators, including information for performance measures. The design allows for more rapid and frequent data reporting (Core studies) and serves as a vehicle for studying more complex issues and topics in greater detail and with increased efficiency (Plus studies).

In fall 2014 and spring 2015, the FACES Core study will assess the school readiness skills of Head Start children, survey their parents, and ask their Head Start teachers to rate children's social and emotional skills. In spring 2015 and again in spring 2017, the number of programs in the FACES Core study sample will increase from the 60 that are used to collect data on children's school readiness outcomes to 180 for the purpose of conducting observations in 720 Head Start classrooms. Program director, center director, and teacher surveys will also be conducted at these time points. FACES Plus studies include additional survey content of policy or programmatic interest, and may involve more programs being sampled. This notice is specific to the data collection activities needed to recruit Head Start programs and centers into FACES. A future notice will provide information about data collection for the Core and Plus studies.

The method of data collection for recruitment of all programs (180 for the FACES Core and up to 50 additional programs for FACES Plus studies) will include telephone conversations with program directors and on-site coordinators who serve as liaisons between the FACES study team and the Head Start centers. These calls will inform program staff about the purpose of the study and will be used to identify the number of centers in each program in order to compile the center sampling frame.

The purpose of this data collection is to support the 2007 reauthorization of the Head Start program (Pub.L. 110–134), which calls for periodic assessments of Head Start's quality and effectiveness.

Respondents: Head Start Program Directors and Staff.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hour per response	Estimated annual burden hours
Telephone script for program directors	230	77	2	1	154
Telephone script for on-site coordinators	230	77	2	.75	116
Total					270

Additional Information: Copies of the proposed collection may be obtained 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. All requests should be identified by the title of the information collection. Email address: *OPREinfocollection@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: 202–395–6974, Attn: Desk Officer for the Administration for Children and Families.

Karl Koerper

OPRE Reports Clearance Officer.

[FR Doc. 2014–02949 Filed 2–11–14; 8:45 am]

BILLING CODE 4184–22–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–0001]

Synergizing Efforts in Standards Development for Cellular Therapies and Regenerative Medicine Products; Public Workshop

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop.

The Food and Drug Administration (FDA), Center for Biologics Evaluation and Research (CBER), is announcing a public workshop entitled ''Synergizing Efforts in Standards Development for Cellular Therapies and Regenerative Medicine Products.'' The purpose of the public workshop is to bring together a broad range of stakeholders to discuss current and future standards development activities involving cellular therapies and regenerative medicine products. This public workshop is being rescheduled due to the government shutdown.

Date and Time: The public workshop will be held on March 31, 2014, from 8:30 a.m. to 4:30 p.m.

Location: The public workshop will be held at the FDA White Oak Campus, 10903 New Hampshire Ave., Bldg. 31 Conference Center, the Great Room (Rm. 1503 A), Silver Spring, MD 20993–0002. Entrance for the public workshop participants (non-FDA employees) is through Building 1 where routine security check procedures will be performed. Please visit the following Web site for location, parking, security, and travel information: *http://www.fda.gov/AboutFDA/WorkingatFDA/BuildingsandFacilities/WhiteOakCampusInformation/ucm241740.htm.*

Contact Person: Sherri Revell, Center for Biologics Evaluation and Research (HFM–49), Food and Drug Administration, 1401 Rockville Pike, Suite 200N, Rockville, MD 20852–1448, 301–827–2000, FAX: 301–827–3079, email: *CBERPublicEvents@fda.hhs.gov* (Subject line: SESDCTRMP Workshop).

Registration: Mail or fax your registration information (including name, title, firm name, address, telephone, and fax numbers) to Sherri Revell (see *Contact Person*) or email to *CBERPublicEvents@fda.hhs.gov* (Subject line: SESDCTRMP Workshop Registration) by March 24, 2014. There is no registration fee for the public workshop. Early registration is recommended because seating is limited. Registration on the day of the public workshop will be provided on a space available basis beginning at 7:30 a.m.

Streaming Webcast of the Public Workshop: For those unable to attend in person, FDA will Webcast the public workshop. To join the Webcast of the public workshop, please go to: *https://collaboration.fda.gov/sesdctrmpworkshop/.* If you have never attended an Adobe Connect meeting before, test your connection at *https://collaboration.fda.gov/common/help/en/support/meeting_test.htm.* Get a quick overview: *http://www.adobe.com/go/connectpro_overview.* Registration is not required for those attending via Adobe Connect.

If you need special accommodations due to a disability, please contact Sherri Revell (see *Contact Person*) at least 7 days in advance.

SUPPLEMENTARY INFORMATION: Standardization efforts concerning the clinical development of cellular therapies and regenerative medicine products have generated a great deal of interest. These efforts include standards development, expert opinion position papers, and professional practice guidelines. However, relatively little is done to coordinate the various existing efforts. In the public workshop, FDA hopes to bring together a broad range of stakeholders of cellular therapies and regenerative medicine products in order to:

- Inform stakeholders about the types of standards and standards organizations that are available currently, the role that the Federal Agencies play in standards development, and the potential role that stakeholders can play in standards development.
- Provide a high-level overview of current standards development activities in the fields of cellular therapy and regenerative medicine and the regulatory application of standards.
- Provide opportunity for discussion of areas of high interest for current or future standards development in the fields of cellular therapy and regenerative medicine and to explore ways to minimize redundancy and maximize collaboration.

We encourage all who have an interest in the development of cellular therapies and regenerative medicine products to attend the public workshop.

This public workshop is being rescheduled due to the government shutdown. It was originally scheduled for October 7, 2013 (see 78 FR 43889, July 22, 2013). Those who registered for the original workshop date must register again for the rescheduled date (see *Registration*).

Transcripts: Please be advised that as soon as possible after a transcript of the public workshop is available, it will be accessible at: *http://www.fda.gov/BiologicsBloodVaccines/NewsEvents/WorkshopsMeetingsConferences/TranscriptsMinutes/default.htm.* Transcripts of the public workshop may also be requested in writing from the Division of Freedom of Information (ELEM–1029), Food and Drug Administration, 12420 Parklawn Dr., Element Bldg., Rockville, MD 20857.

Dated: February 6, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2014–03015 Filed 2–11–14; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–0001]

Anti-Infective Drugs 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: Anti-Infective Drugs 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 March 31, 2014, from 8 a.m. to 5 p.m.

Location: FDA White Oak Campus, 10903 New Hampshire Ave., Bldg. 31 Conference Center, the Great Room (rm. 1503), Silver Spring, MD 20993–0002. Information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: *http://www.fda.gov/AdvisoryCommittees/default.htm;* under