self-sufficiency services for parents, parent skills training, and high-quality child care for children in low-income families in Kansas and Missouri.

The purpose of the current document is to request public comment on the 36-month participant survey in Philadelphia. The research team plans to collect participant-reported surveys assessing participants' employment, education and economic outcomes, participation in employment and training services, receipt of benefits and services such as food stamps and mental health services, housing and household information, health and health care coverage, child care, and child outcomes.

The follow-up survey at the 36-month follow-up in Philadelphia will be used for the following purposes: To study the extent to which pre-employment services and transitional employment affect employment, earnings, income, and welfare dependence of low-income TANF recipients; to examine the impacts of these services on participants' health, receipt of benefits such as food stamps, Medicaid, and child-care subsidies, and participation in services such as substance abuse treatment and mental health services; and to collect data on a wider range of outcomes measures than is available through welfare, Medicaid, Food Stamps, Social Security, and Unemployment Insurance records.

The 36-month data collection effort draws heavily from the 15-month survey conducted in this site. Materials for the 15-month data collection effort were previously submitted to OMB and were approved (OMB Control No. 0970–0276).

Respondents: TANF recipients without a high school diploma and/or recipients who have received TANF for at least 12 months.

The fielded sample of the 36-month data collection effort will be all 1,944 participants in the two program groups and the control group of the HtE project in Philadelphia. The burden estimates below assume an 80 percent response rate of the fielded sample.

The annual burden estimates are detailed below, and the substantive content of each component will be detailed in the supporting statement attached to the forthcoming 30-day notice.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Philadelphia 36-month participant survey	1,555	1	0.75	1,166

Estimated Total Annual Burden Hours: 1,166.

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 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: June 13, 2007.

Brendan Kelly,

OPRE Reports Clearance Officer.

[FR Doc. 07–2998 Filed 6–18–07; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: Request for State Data Needed to Determine the Amount of a Tribal Family Assistance Grant.

OMB No.: 0970–0173.

Description: 42 U.S.C. 612 (Section 412 of the Social Security Act) gives federally recognized Indian Tribes the opportunity to apply to operate a Tribal Temporary Assistance for Needy Families (TANF) program. The Act specifies that the Secretary shall use State-submitted data to determine the amount of the grant to the Tribe. This form (letter) is used to request those data from the States. ACF is proposing to extend this information collection without change.

Respondents: States that have Indian Tribes applying to operate a TANIF program.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Request for State Data Needed to Determine the Amount of a Tribal Family Assistance Grant	15	1	42	630

Estimated Total Annual Burden Hours: 630.

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: 202–395–6974, Attn: Desk Officer for the Administration for Children and Families.

Dated: June 14, 2007.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 07–3004 Filed 6–18–07; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Cellular, Tissue, and Gene Therapies 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). At least one portion of the meeting will be closed to the public.

Name of Committee: Cellular, Tissue, and Gene Therapies 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 by teleconference on July 26, 2007 from 11 a.m. to approximately 3:15 p.m.

Location: National Institutes of Health, Bldg. 29B, Conference rms. A/B, 9000 Rockville Pike, Bethesda, MD. This meeting will be held by teleconference. The public is welcome to attend the meeting at the specified location. A speakerphone will be provided at the specified location for public participation in the meeting. Important information about transportation and directions to the NIH campus, parking, and security procedures is available on the Internet at *http://www.nih.gov/about/visitor/index.htm*. Visitors must show two forms of identification, one of which must be a government-issued photo identification such as a Federal employee badge, driver's license, passport, green card, etc. If you are planning to drive to and park on the NIH campus, you must enter at the South Dr. entrance of the campus which is located on Wisconsin Ave. (the Medical Center Metro entrance), and allow extra time for vehicle inspection. Detailed information about security procedures is located at *http://www.nih.gov/about/visitorsecurity.htm*. Due to the limited available parking, visitors are encouraged to use public transportation.

Contact Person: Gail Dapolito or Danielle Cubbage, Center for Biologics Evaluation and Research, Food and Drug Administration, 1401 Rockville Pike, Rockville, MD, 20852, 301–827–0314, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 3014512389. 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: On July 26, 2007, the committee will meet in open session to hear updates of research programs in: (1) The Division of Therapeutic Proteins and the Division of Monoclonal Antibodies, Office of Biotechnology Products, Center for Drug Evaluation and Research, FDA and (2) the Division of Cellular and Gene Therapies, Office of Cellular, Tissue, and Gene Therapies, Center for Biologics Evaluation and Research, FDA.

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/ohrms/dockets/ac/acmenu.htm*, click on the year 2007 and scroll down to the appropriate advisory committee link.

Procedure: On July 26, 2007, from 11 a.m. to approximately 2 p.m., the meeting is open to the public. 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 July 12, 2007. 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 statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before July 18, 2007. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by July 19, 2007.

Closed Committee Deliberations: On July 26, 2007, from approximately 2:15 p.m. to 3:30 p.m., the meeting will be closed to permit discussion where disclosure would constitute a clearly unwarranted invasion of personal privacy (5 U.S.C. 552b(c)(6)). The committee will discuss reports of intramural research programs and make recommendations regarding personnel staffing decisions.

Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Gail Dapolito at least 7 days in advance of the meeting.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: June 11, 2007.

Randall W. Lutter,

Associate Commissioner for Policy and Planning.

[FR Doc. E7–11728 Filed 6–18–07; 8:45 am]

BILLING CODE 4160–01–S