The response burden for service providers is estimated as:

Component	Number of respondents	Responses per provider	Total responses	Hours per response	Total burden hours
Provider Report	2,080*	1	2,080*	2.30	4,784

Component	Electronic data system	Number of respondents	Responses per provider	Total responses	Hours per response	Total burden hours
Client Report	No	56	1	56	106.25	5,950
	Yes	1,822	1	1,822	3.75	6,832.5
	Subtotal	**1,878		**1,878		12,782.5

* All providers, including providers of administrative support services and direct client services.
** Providers of direct client services only.

Total Burden Hours: 17,974.5

E-mail comments to *paperwork@hrsa.gov* or mail comments to the HRSA Reports Clearance Officer, Room 10–33, Parklawn Building, 5600 Fishers Lane, Rockville, Maryland, 20857. Written comments should be received within 60-days of this notice. Information can also be accessed at *http://datasupport.hab.hrsa.gov/*.

Dated: September 14, 2010.

Sahira Rafiullah,

Director, Division of Policy and Information Coordination.

[FR Doc. 2010–23416 Filed 9–17–10; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Request for State Data Needed to Determine 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 TANF program.

ANNUAL BURDEN ESTIMATES

Information collection	Number of respondents	Number of responses per respondent	Average burden per response	Total burden hours
Request for State Data Needed to Determine the Amount of Tribal Family Assistance Grant	4	1	42	168

Total Estimated Burden: 168 hours.

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: September 13, 2010.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2010–23319 Filed 9–17–10; 8:45 am]

BILLING CODE M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0001]

Risk Communication 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: Risk Communication 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 November 8, 2010, from 8 a.m. to 5 p.m. and November 9, 2010, from 8 a.m. to 2 p.m.

Location: FDA White Oak Campus, Building 31 Conference Center, Great Room, 10903 New Hampshire Ave., Silver Spring, MD 20993. Please note visitors can park in the southwest garage near Building 31 or the northwest parking lot near Building 22 (for a campus map, see *http://www.fda.gov/downloads/AboutFDA/WorkingatFDA/BuildingsandFacilities/WhiteOakCampusInformation/UCM194893.pdf*). Visitors to the White Oak Campus must have a valid driver's license or other picture ID, and must enter through Building 1.

Contact Person: Lee L. Zwanziger, Office of Policy, Planning and Preparedness, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, rm. 3278, Silver Spring, MD, 20993, 301–796–9151, FAX: 301–847–8611, e-mail: *RCAC@fda.hhs.gov*, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 8732112560. 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 November 8 and 9, 2010, the Committee will hear and discuss developments in FDA's ongoing communications programs, such as FDA's Strategic Plan for Risk Communication, FDA's Transparency Initiative, and the challenges of effectively communicating with patients and caregivers about appropriate use of medical devices when a patient is prescribed a medical device for home use.

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 October 29, 2010. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. on November 8, 2010, and 10:30 to 11:30 a.m. on November 9, 2010. 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 October 21, 2010. 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 October 22, 2010.

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 Lee Zwanziger at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at *http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm* for procedures on public conduct during advisory committee meetings.

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

Dated: September 14, 2010.

Jill Hartzler Warner,

Acting Associate Commissioner for Special Medical Programs.

[FR Doc. 2010–23368 Filed 9–17–10; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Environmental Health Sciences; Notice of Meeting

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

The meeting will be open to the public as indicated below, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the meeting.

The meeting will be closed to the public as indicated below in accordance with the provisions set forth in section 552b(c)(6), Title 5 U.S.C., as amended for the review, discussion, and evaluation of individual intramural programs and projects conducted by the NATIONAL INSTITUTE OF ENVIRONMENTAL HEALTH SCIENCES, including consideration of personnel qualifications and performance, and the competence of individual investigators, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Board of Scientific Counselors, NIEHS.

Date: October 17–19, 2010.

Closed: October 17, 2010, 7 p.m. to 10 p.m

Agenda: To review and evaluate programmatic and personnel issues.

Place: Doubletree Guest Suites, 2515 Meridian Parkway, Research Triangle Park, NC 27713.

Open: October 18, 2010, 8:30 a.m. to 11:50 a.m.

Agenda: An overview of the organization and research in the Laboratory of Reproductive and Developmental Toxicology.

Place: Nat. Inst. of Environmental Health Sciences, Building 101, Rodbell Auditorium, 111 T. W. Alexander Drive, Conference Rooms 101 A, B, and C, Research Triangle Park, NC 27709.

Closed: October 18, 2010, 11:50 a.m. to 12:35 p.m.

Agenda: To review and evaluate programmatic and personnel issues.

Place: Nat. Inst. of Environmental Health Sciences, Building 101, Rodbell Auditorium, 111 T. W. Alexander Drive, Conference Rooms 101 A, B, and C, Research Triangle Park, NC 27709.

Open: October 18, 2010, 1:30 p.m. to 2:45 p.m.

Agenda: An overview of the organization and research in the Laboratory of Reproductive and Developmental Toxicology.