notify interested persons regarding their request to speak by August 8, 2008.

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 L. 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/oc/advisory/default.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: July 17, 2008.

Randall W. Lutter,

Deputy Commissioner for Policy.

[FR Doc. E8–17304 Filed 7–28–08; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget (OMB), in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to OMB for review, e-mail *paperwork@hrsa.gov* or call the HRSA Reports Clearance Office on (301) 443–1129.

The following request has been submitted to the Office of Management and Budget for review under the Paperwork Reduction Act of 1995:

Proposed Project: Sickle Cell Disease Treatment Demonstration Program (SCDTDP), Health Resources and Services Administration (HRSA): NEW

In 2004 Congress enacted and the President signed into law Public Law 108–357, the American Jobs Creation Act of 2004. Section 712 of Public Law 108–357 authorized a demonstration program for the prevention and treatment of Sickle Cell Disease. The legislation was enacted to (1) create an optional medical assistance program for individuals with Sickle Cell Diseases for treatment and education, genetic counseling and other services to prevent mortality and decrease morbidity from Sickle Cell Disease, and (2) create a demonstration program, the SCDTDP, under HRSA. The SCDTDP provides grants to federally-qualified and nonprofit health care providers to establish geographically distributed regional networks that will work with comprehensive Sickle Cell Disease centers and community-based support organizations to provide coordinated, comprehensive, culturally competent, and family-centered care to families with Sickle Cell Disease. In fiscal year 2006, HRSA awarded four, 4-year grants to the Illinois Sickle Cell Association Network, Alabama Network for Sickle Cell Care, Access, Prevention, and Education, Carolina Partnership for Sickle Cell Treatment Continuum of Care, and the Cincinnati Sickle Cell Network.

Under the authorizing legislation, a National Coordinating Center (NCC) was established to (1) collect, coordinate, monitor, and distribute data, best practices and findings regarding the activities of the demonstration program, (2) identify a model protocol for eligible entities with respect to the prevention and treatment of Sickle Cell Disease, (3) identify educational materials regarding the prevention and treatment of Sickle Cell Disease, and (4) prepare a final report on the efficacy of the demonstration program based on evaluation findings.

As part of the evaluation, pre and post utilization and satisfaction data and quality of life assessments will be collected from the demonstration clients during various phases of their participation. These data will be collected through medical record abstractions and self-report using hard copy questionnaires and submitted to the NCC for processing and analysis.

The total burden estimate per participant is shown below:

Type of respondent	Form name	Number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
Sickle Cell Disease clients or caregivers.	Utilization Questionnaire (pre-demonstration).	400	1	400	.75	300
Sickle Cell Disease clients or caregivers.	Utilization Questionnaire (post demonstration).	400	1	400	.50	200
Sickle Cell Disease clients or caregivers.	SF–36 Health Survey for adults over 18 years of age.	280	2	560	.25	140
Parents of Sickle Cell Disease clients.	PedsQL for parents	120	2	240	.25	60
Sickle Cell Disease clients age 18 and younger.	PedsQL for children and adolescents.	100	2	200	.25	50
Sickle Cell Disease clients or caregivers.	The Medical Home Family Index (Health Care Satisfaction).	400	2	800	.25	200
Total		500		2,600		950

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to the desk officer for HRSA, either by e-mail to *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974. Please direct all correspondence to the ''attention of the desk officer for HRSA.''

Dated: July 23, 2008.

Alexandra Huttinger,

Director, Division of Policy Review and Coordination.

[FR Doc. E8–17354 Filed 7–28–08; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of the following meetings.

The meetings 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: Center for Scientific Review Special Emphasis Panel Member Conflict: Health Risks, Interventions and Outcomes.

Date: August 5, 2008.

Time: 11 a.m. to 3 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892. (Virtual Meeting)

Contact Person: Michael Micklin, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3136, MSC 7759, Bethesda, MD 20892, (301) 435–1258, *micklinm@csr.nih.gov*.

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

Name of Committee: Center for Scientific Review Special Emphasis Panel, Microcirculation and Renovascular Hypertension.

Date: August 21, 2008.

Time: 2 p.m. to 4 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892. (Telephone Conference Call)

Contact Person: Olga A. Tjurmina, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4030B, MSC 7814, Bethesda, MD 20892, (301) 451–1375, *ot3d@nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel, Cardiac Fibrillation and Defibrillation.

Date: September 5, 2008.

Time: 2 p.m. to 4 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892. (Telephone Conference Call)

Contact Person: Maqsood A. Wani, PhD, DVM, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 2114, MSC 7814, Bethesda, MD 20892, 301–435–2270, *wanimaqs@csr.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel, NIH Rapid Access to Interventional Development Review.

Date: September 17–18, 2008.

Time: 8 a.m. to 8 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892. (Virtual Meeting)

Contact Person: James J. Li, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5148, MSC 7849, Bethesda, MD 20892, 301–435–2417, *lijames@csr.nih.gov*.

Name of Committee: Oncological Sciences Integrated Review Group, Developmental Therapeutics Study Section.

Date: September 18–19, 2008.

Time: 8 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: Doubletree Hotel, 1515 Rhode Island Avenue, NW., Washington, DC 20005.

Contact Person: Sharon K. Gubanich, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6214, MSC 7804,Bethesda, MD 20892, (301) 435–1767, *gubanics@csr.nih.gov*.

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393–93.396, 93.837–93.844, 93.846–93.878, 93.892, 93.893, National Institutes of Health, HHS)

Dated: July 21, 2008.

Jennifer Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. E8–17166 Filed 7–28–08; 8:45 am]

BILLING CODE 4140–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Alcohol Abuse and Alcoholism; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), 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 on Alcohol Abuse and AlcoholismSpecial Emphasis Panel; Applications in Response to PAR–07–157.

Date: August 19, 2008.

Time: 1 p.m. to 3 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institute on Alcohol Abuse and Alcoholism, 5635 Fishers Lane, Room 3146, Rockville, MD 20852, (Telephone Conference Call).

Contact Person: Katrina L Foster, PhD, Scientific Review Officer, National Inst on Alcohol Abuse & Alcoholism, National Institutes of Health, 5635 Fishers Lane, Rm. 3042,Rockville, MD 20852, 301–443–4032, *Katrina@mali.ruh.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.271, Alcohol Research Career Development Awards for Scientists and Clinicians; 93.272, Alcohol National Research Service Awards for Research Training; 93.273, Alcohol Research Programs; 93.891, Alcohol Research Center Grants, National Institutes of Health, HHS)

Dated: July 18, 2008.

Jennifer Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. E8–17165 Filed 7–28–08; 8:45 am]

BILLING CODE 4140–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Alcohol Abuse and Alcoholism; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), 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,