

Name of Committee: National Center on Minority Health and Health Disparities.

Date: May 10–12, 2010.

Time: 6 p.m. to 5 p.m.

Agenda: To review and evaluate grant applications and or proposals.

Place: Embassy Suites at the Chevy Chase Pavilion, 4300 Military Road, NW., Washington, DC 20015.

Contact Person: Prabha L. Atreya, PhD, Chief, Office of Scientific Review, National Center on Minority Health and Health Disparities, 6707 Democracy Boulevard, Suite 800, Bethesda, MD 20892, (301) 594–8696, atreyapr@mail.nih.gov.

Dated: April 20, 2010.

Jennifer Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010–9638 Filed 4–23–10; 8:45 am]

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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. App.), 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; OBT IRG Member Conflict.

Date: May 17, 2010.

Time: 3 p.m. to 5 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: Angela Y. Ng, MBA, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6200, MSC 7804 (For courier delivery, use MD 20817), Bethesda, MD 20892, 301–435–1715, nga@csr.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Healthcare Delivery and Methodologies Competitive Supplements.

Date: May 18, 2010.

Time: 11:30 a.m. to 1 p.m.

Agenda: To review and evaluate grant applications.

Place: Sheraton Sand Key Hotel, 1160 Gulf Boulevard, Clearwater Beach, FL 33767.

Contact Person: Jacinta Bronte-Tinkew, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3164, MSC 7770, Bethesda, MD 20892, (301) 435–1503, brontetinkewjm@csr.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: Cancer Genetics and Epidemiology.

Date: May 19, 2010.

Time: 12 p.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: Tata Communications, 2355 Dulles Corner Boulevard, 7th Floor, Herndon, VA 20171 (Virtual Meeting).

Contact Person: Fungai Chanetsa, MPH, PhD., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3135, MSC 7770, Bethesda, MD 20892, 301–408–9436, fungai.chanetsa@nih.hhs.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: April 20, 2010.

Jennifer Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010–9652 Filed 4–23–10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0204]

Infusion Pumps; Public Meeting; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public meeting; request for comments.

SUMMARY: The Food and Drug Administration (FDA) is announcing a public meeting regarding external infusion pumps. The purpose of the meeting is to inform the public about current problems associated with external infusion pump use, to help the agency identify quality assurance strategies to mitigate these problems, and to solicit comments and input regarding how to bring more effective external infusion pumps to market. Elsewhere in this issue of the **Federal Register**, FDA is announcing the availability of a draft guidance document entitled “Total Product Life Cycle: Infusion Pump—Premarket Notification (510(k)) Submissions.”

Date and Time: The public meeting will be held on May 25 and 26, 2010,

from 8 a.m. to 5 p.m. Persons interested in attending the meeting must register by 5 p.m. on May 18, 2010.

Location: The public meeting will be held at the Hilton Silver Spring hotel, 8727 Colesville Rd., Silver Spring, MD 20910. Seating is limited and available only on a first-come, first-served basis.

Contact Person: Victoria Wagman, Food and Drug Administration, Center for Devices and Radiological Health (CDRH), 10903 New Hampshire Ave., Bldg. 66, rm. 5449, Silver Spring, MD 20993–0002, 301–796–6851, e-mail: victoria.wagman@fda.hhs.gov.

Registration: Register online for webinar or onsite attendance at <http://www.fda.gov/MedicalDevices/NewsEvents/WorkshopsConferences/ucm203299.htm> (select the appropriate meeting from the list). Please provide complete contact information for each attendee, including name, title, affiliation, address, e-mail, and telephone number. Registration requests should be received by May 18, 2010. For those without Internet access, please call 301–796–6861 to register.

Registration is free and will be on a first-come, first-served basis. Early registration is recommended because seating is limited and therefore FDA may limit the number of participants from each organization. If time and space permits, onsite registration on the day of the public meeting will be provided beginning at 7 a.m. Persons needing a sign language interpreter or other special accommodations should notify Victoria Wagman (see *Contact Person*) at least 7 days in advance. Additional information is also available at <http://www.fda.gov/MedicalDevices/NewsEvents/WorkshopsConferences/ucm181140.htm>.

Comments: Regardless of attendance at the public meeting, interested persons may submit written or electronic comments. Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to <http://www.regulations.gov>. Submit a single copy of electronic comments or two paper copies of any mailed comments, except that individuals may submit one paper copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Please also indicate the specific question(s) addressed. (See section II of this document.) Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

SUPPLEMENTARY INFORMATION: