DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Program Announcement 04199]

Rapid Expansion of HIV/AIDS Activities by National Ivorian Nongovernmental Organizations and Associations Serving Highly Vulnerable Populations in Cote d'Ivoire Under the President's Emergency Plan for AIDS Relief; Amendment

A notice announcing the availability of fiscal year (FY) 2004 funds for cooperative agreements for Rapid Expansion of HIV/AIDS Activities by National Ivorian Nongovernmental Organizations and Associations Serving Highly Vulnerable Populations in Cote d'Ivoire Under the President's Emergency Plan for AIDS Relief was published in the **Federal Register** on June 1, 2004, volume 69, Number 105, pages 30922–30927.

The notice is amended as follows:

On page 30923, under ''Activities: Awardee activities for this program are as follows:'' please include:

For Year 1, the grantee will focus on building on existing interventions with transactional sex workers, their partners, and other vulnerable women at sites including, but not limited to: the Clinique de Confiance, Cote d'Ivoire Properite, APROSAM.''

Also on page 30923, ''III.1. Eligible Applicants'', please change the first sentence to read: ''Applications may be submitted by international nongovernmental organizations, including faith-based organizations, that have experience in: designing and implementing HIV/AIDS activities for Highly Vulnerable Populations in general, and transactional sex workers in particular, in Africa; capacity building for African local nongovernmental organizations and associations in developing countries (including resource mobilization and administration of funds); and understanding complexities and challenges of designing and implementing activities for HVP.''

Dated: June 25, 2004.

Alan Kotch,

Acting Director, Procurement and Grants Office, Centers for Disease Control and Prevention.

[FR Doc. 04–14938 Filed 6–30–04; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Circulatory System Devices Panel of the Medical Devices 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: Circulatory System Devices Panel of the Medical Devices 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 July 28, 2004, from 9 a.m. to 5 p.m., and on July 29, 2004, from 9 a.m. to 5 p.m.

Location: Holiday Inn, Grand Ballroom, Two Montgomery Village Ave., Gaithersburg, MD.

Contact Person: Geretta Wood, Center for Devices and Radiological Health (HFZ–450), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–443–8320, ext. 143, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 3014512625. Please call the Information Line for up-to-date information on this meeting.

Agenda: On July 28, 2004, the committee will discuss, make recommendations, and vote on a premarket approval application supplement for a cardiac resynchronization device. On July 29, 2004, the committee will hear a presentation on Adverse Event Reports for Automatic External Defibrillators from 1996 to 2003.

The committee will also discuss and make recommendations on a premarket notification (510(k)) submission for an over-the-counter automated external defibrillator. Background information for the topics, including the agenda and questions for the committee, will be available to the public 1 business day before the meeting on the Internet at *http://www.fda.gov/cdrh/panelmtg.html*. Material for the July 28 session will be posted on July 27, 2004; material for the July 29 session will be posted on July 28, 2004.

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 by July 14, 2004. On both days, oral presentations from the public will be scheduled for approximately 30 minutes at the beginning of committee deliberations and for approximately 30 minutes near the end of the deliberations. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before July 14, 2004, 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.

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 AnnMarie Williams, Conference Management Staff, at 301–594–1283, ext. 113, 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 24, 2004.

William K. Hubbard,

Associate Commissioner for Policy and Planning.

[FR Doc. 04–14947 Filed 6–30–04; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Cancer Institute; 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/or contract proposals and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant