20993, 301–796–6313, e-mail: James.Swink@fda.hhs.gov, 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. 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 March 18, 2010, the committee will discuss, make recommendations, and vote on the premarket approval application (PMA) for the Cardiac Resynchronization Therapy Defibrillators (CRT–Ds) sponsored by Boston Scientific. The sponsor is seeking expanded indications for the their CRT–Ds to include patients with low left ventricular ejection fraction (<30%) and wide QRS (≥130 ms) who are NYHA Class II (ischemic or non-ischemic etiology) or NYHA Class I (ischemic etiology).

On March 19, 2010, the committee will discuss, make recommendations and vote on a PMA for the REVO MRI Pacing System sponsored by Medtronic. The REVO MRI Pacing System is a pacemaker (with a standard pacing indication) that has been specifically designed to be safe for the MRI environment under certain MR scanning conditions.

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. 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 March 11, 2010. Oral presentations from the public will be scheduled immediately following lunch. Those desiring to make 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 March 3, 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 March 4, 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 AnnMarie Williams at 301–796–5966 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).


Jill Hartzler Warner, Acting Associate Commissioner for Special Medical Programs.

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

**Board of Scientific Counselors, Coordinating Center for Infectious Diseases, (BSC, CCID)**

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC), announces the following meeting of the aforementioned committee:

**Time and Date:** 12 p.m.–2 p.m., April 7, 2010 (Closed).

**Place:** Teleconference.

**Status:** The meeting will be closed to the public in accordance with provisions set forth in Section 552b(c)(4) and (6), Title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92–463.

**Matters To Be Discussed:** The meeting will include the initial review, discussion, and evaluation of applications received in response to “Strengthening Global Human-Animal Interface Activities for Avian Influenza and other Zoonotic Diseases, FOA CK10–001.”

**Contact Person for More Information:**
Gregory Anderson, M.S., M.P.H., Scientific Review Officer, CDC, 1600 Clifton Road, NE., Mailstop E60, Atlanta, GA 30333, Telephone: (404) 498–2293.

The Director, Management Analysis and Services Office, has been delegated the authority to sign Federal Register notices pertaining to announcements of meetings and other committee management activities, for both CDC and the Agency for Toxic Substances and Disease Registry.


Andre Tyler, Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

**Purpose:** The BSC, CCID shall advise the Secretary, HHS, and the Director, CDC concerning strategies and goals for the programs and research within the national centers; will administer and oversee peer review of scientific programs; and monitor the overall strategic direction and focus of the national centers.

**Matters To Be Discussed:** Agenda items will include:
1. Update from Dr. Khabbaz.
2. Update on H1N1 response.