Oak Campus.” Please note that visitors to the White Oak Campus must enter through Building 1.

Contact Person: Martha Monser, Office of the Chief Scientist, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, rm. 4286, Silver Spring, MD 20993, 301–796–4627, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area). 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 at http://www.fda.gov/AdvisoryCommittees/default.htm and scroll down to the appropriate advisory committee meeting link, or call the advisory committee information line to learn about possible modifications before coming to the meeting.

Agenda: On February 27, 2013, the Science Board will be provided with updates and/or draft reports from the Center for Devices and Radiological Health Research Review subcommittee and the Global Health subcommittee. A progress update will be presented regarding the recently established Center for Biologics Evaluation and Review Post-Marketing Safety Review subcommittee. An update on the plan to establish a new subcommittee to evaluate the Agency’s continuing work to address the challenges identified in Science Board’s 2007 “Science and Mission at Risk” Report will be presented. Plans will also be presented and a proposed charge reviewed for a second subcommittee to assess similar issues with respect to information technology including evaluation of scientific data utilization, data liberation and innovation. Overviews of genomics activities at the Centers for Food Safety and Applied Nutrition and Veterinary Medicine will be presented, along with plans for an Agency-wide working group to address cross-cutting genomics activities. Finally, recipients of the FY2012 Scientific Achievement Awards (selected by the Science Board) will provide overviews of the activities for which the awards were given.

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 Wednesday, February 20, 2013. Oral presentations from the public will be scheduled between approximately 4:30 p.m. and 5 p.m. Those individuals interested in making 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 Tuesday, February 12, 2013. 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 Wednesday, February 13, 2013.

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 disabilities or special needs, if you require special accommodations due to a disability, please contact Martha Monser 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: January 24, 2013.

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

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