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DEPARTMENT OF HEALTH AND HUMAN SERVICES

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Food and Drug Administration

National Mammography Quality Assurance 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: National Mammography Quality Assurance 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 September 26 and 27, 2005, from 9 a.m. to 6 p.m.

Location: Holiday Inn, Walker/Whetstone Rooms, Two Montgomery Village Ave., Gaithersburg, MD.

Contact Person: Charles Finder, Center for Devices and Radiological Health (HFZ-240), Food and Drug Administration, 1350 Piccard Dr., Rockville, MD 20850, 301-594-3332, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 3014512397.

Please call the Information Line for up-to-date information on this meeting.

Agenda: The committee will discuss the following issues:

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- (1) Regulatory and nonregulatory mechanisms to enhance mammography quality while reducing the regulatory and inspection burden on facilities;
- (2) Recommendations made by the Institute of Medicine regarding the current Mammography Quality Standards Act (MQSA) program, interventional mammography, and nonmammographic breast imaging procedures; and
- (3) All relevant guidance documents issued since the last meeting.

The committee will also receive updates on recently approved alternative standards, voluntary stereotactic accreditation programs, and the radiological health program. MQSA regulations and guidance documents are available to the public on the Internet at <http://www.fda.gov/cdrh/mammography>.

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 September 5, 2005. Oral presentations from the public will be scheduled between approximately 9:30 a.m. and 10:30 a.m. on both days. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before September 5, 2005, 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.

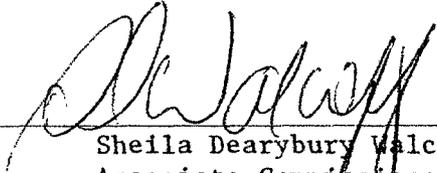
Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

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 Shirley Meeks at 240-276-0450, ext. 105, 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: 7/27/05
July 27, 2005.



Sheila Dearybury Walcoff,
Associate Commissioner for External Relations.

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