Marianne Augustine at (202) 690–7102 by 5 p.m. E.D.T., September 19, 2003. Space is limited for all sessions. Written comments from the public will be accepted; opportunities to present oral comments may be provided at future meetings. Please call Marianne Augustine by 5 p.m. E.D.T., September 12, 2003, should you require a sign language interpreter. Documents pertaining to Committee deliberations will be available for public inspection and copying in Room 738-G, 200 Independence Avenue, SW., Washington, DC 20201 on the day before the meeting and following the meeting. Please call (202) 690–7102 to schedule an appointment to view the documents.

Written Comment: By this notice, the Committee is soliciting written comments, views, information and data pertinent to review of the Dietary Guidelines for Americans. Written comments are welcome throughout the Committee's deliberations. To be considered for the first meeting, they must be received by 5 p.m. E.D.T. on September 16, 2003. Comments should be sent to

dietaryguidelines@osophs.dhhs.gov or to Kathryn McMurry, HHS Office of Disease Prevention and Health Promotion, Room 738–G, 200 Independence Avenue, SW., Washington, DC 20201.

Dated: August 26, 2003.

Carter Blakey,

Acting Deputy Assistant Secretary for Health, U.S. Department of Health and Human Services.

Dated: August 27, 2003.

Eric J. Hentges,

Executive Director, Center for Nutrition Policy and Promotion, U.S. Department of Agriculture.

Dated: August 28, 2003.

Edward Knipling,

Acting Administrator, Agricultural Research Service, U.S. Department of Agriculture. [FR Doc. 03–22480 Filed 9–3–03; 8:45 am] BILLING CODE 4150–32–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Notice of Meetings

In accordance with section 10(d) of the Federal Advisory Committee Act as amended (5 U.S.C., Appendix 2), the Agency for Healthcare Research and Quality (AHRQ) announces meetings of scientific peer review groups. The subcommittees listed below are part of the Agency's Health Services Research Initial Review Group Committee.

The subcommittee meetings will be closed to the public in accordance with the Federal Advisory Committee Act, section 10(d) of 5 U.S.C., Appendix 2 and 5 U.S.C. 552b(c)(6). Grant applications are to be reviewed and discussed at these meetings. These discussions are likely to involve information concerning individuals associated with the applications, including assessments of their personal qualifications to conduct their proposed projects. This information is exempt from mandatory disclosure under the above-cited statutes.

1. *Name of Subcommittee:* Health Care Research Training.

Date: September 25–26, 2003 (Open from 8 a.m. to 8:15 a.m. on September 25 and closed for remainder of the meeting).

Place: AHRQ Conference Center, John M. Eisenberg Bldg, Rockville, Maryland 20850.

2. *Name of Subcommittee:* Health Research Dissemination and Implementation.

Date: October 23–24, 2003 (Open from 8 a.m. to 8:15 a.m. on October 23 and closed for remainder of the meeting).

Place: AHRQ Conference Center, John M. Eisenberg Bldg, Rockville, Maryland 20850. 3. *Name of Subcommittee:* Health Care

Quality and Effectiveness Research.

Date: October 23–24, 2003 (Open from 8 a.m. to 8:15 a.m. on October 23 and closed for remainder of the meeting).

Place: AHRQ Conference Center, John M. Eisenberg Bldg, Rockville, Maryland 20850.

4. Name of Subcommittee: Health Systems Research.

Date: October 27–28, 2003 (Open from 8 a.m. to 8:15 a.m. on October 27 and closed for remainder of the meeting).

Place: AHRQ Conference Center, John M. Eisenberg Bldg, Rockville, Maryland 20850. 5. Name of Subcommittee: Health Care

Technology and Decision Sciences.

Date: October 30–31, 2003 (Open from 8 a.m. to 8:15 a.m. on October 30 and closed for remainder of the meeting).

Place: AHRQ Conference Center, John M. Eisenberg Bldg, Rockville, Maryland 20850.

Contact Person: Anyone wishing to obtain a roster of members, agenda or minutes of the nonconfidential portions of the meetings should contact Mrs. Bonnie Campbell, Committee Management Officer, Office of Extramural Research Review, Education and Priority Populations, AHRQ, 540 Gaither Road, Rockville, Maryland 20850, Telephone (301) 427–1554.

Agenda items for these meetings are subject to change as priorities dictate.

Dated: August 26, 2003.

Carolyn M. Clancy, Director.

[FR Doc. 03–22476 Filed 9–3–03; 8:45 am] BILLING CODE 4160–90–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2003N-0376]

Medical Devices: Mammography Quality Standards Act of 1992 and Subsequent Mammography Quality Standards Reauthorization Act and Amendments; Inspection Fees

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the new fees the agency will assess for inspections of mammography facilities starting October 1, 2003. The Mammography Quality Standards Act of 1992 (the MQSA) requires FDA to assess and collect fees from mammography facilities to cover the costs of annual inspections required by the MQSA. Because these costs have increased since the last increase on February 13, 1998, FDA is raising the fees accordingly. This document explains which facilities are subject to payment of inspection fees, provides information on the costs included in developing inspection fees, and provides information on the inspection billing and collection processes. This is only the second increase in inspection fees under the MQSA since the initial fee was established in 1995.

DATES: Effective October 1, 2003, for all inspections conducted under section 354(g) of the Public Health Service Act (PHS Act) (42 U.S.C. 263b(g)). Submit written comments by October 1, 2003.

ADDRESSES: Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20857. Submit electronic comments to *http:// www.fda.gov/dockets/ecomments*. Identify comments with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: John L. McCrohan, Center for Devices and Radiological Health (HFZ–240), Food and Drug Administration, 1350 Piccard Dr., Rockville, MD 20850, 301–594–3332, FAX: 301–594–3306.

SUPPLEMENTARY INFORMATION:

I. Background

The MQSA requires all mammography facilities, other than facilities of the Department of Veterans Affairs, to be accredited by an approved accreditation body and certified by the