Exposure Draft is hereby postponed from October 1995 to December 1995, actual date to be announced later in the **Federal Register**.

FOR FURTHER INFORMATION CONTACT:

Ronald S. Young, Executive Staff Director, 750 First ST., NE., Room 1001, Washington, DC 20002, or call (202) 512–7350.

Authority: Federal Advisory Committee Act. Pub. L. No. 92–463, Section 10(a)(2), 86 Stat. 770, 774 (1972) (current version at 5 U.S.C. app. section 10(a)(2) (1988); 41 CFR 101–6.1015 (1990).

Dated: October 5, 1995.

Ronald S. Young,

Executive Director.

[FR Doc. 95–25195 Filed 10–10–95; 8:45 am]

BILLING CODE 1610–01–M

Federal Accounting Standards Advisory Board

AGENCY: General Accounting Office.

ACTION: Notice of monthly meeting.

SUMMARY: Pursuant to section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. No. 92–463), as amended, notice is hereby given that the regular monthly meeting of the Federal Accounting Standards Advisory Board will be held on Thursday, October 19 from 9 a.m. to 4 p.m., continuing on Thursday, October 26, and concluding on Friday, October 27, 1995 at noon in room 7C13 of the General Accounting Office, 441 G St., NW., Washington, DC.

The purpose of the meeting is to discuss issues arising from the September 20 public hearing on *Accounting for Revenue and Other Financing Sources* exposure draft and to discuss any other issues related to the exposure draft.

Any interested person may attend the meeting as an observer. Board discussions and reviews are open to the public.

FOR FURTHER INFORMATION CONTACT:

Ronald S. Young, Executive Staff Director, 750 First St., NE., Room 1001, Washington, DC 20002, or call (202) 512–7350.

Authority: Federal Advisory Committee Act. Pub. L. No. 92–463, Section 10(a)(2), 86 Stat. 770, 774 (1972) (current version at 5 U.S.C. app. section 10(a)(2) (1988); 41 CFR 101–6.1015) (1990).

Dated: October 5, 1995.

Ronald S. Young,

Executive Director.

[FR Doc. 95–25194 Filed 10–10–95; 8:45 am]

BILLING CODE 1610–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Advisory Committees; Filing of Annual Reports

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that, as required by the Federal Advisory Committee Act, the agency has filed with the Library of Congress the annual reports of those FDA advisory committees that held closed meetings during fiscal year 1994. FDA apologizes for the lateness in the filing of these reports due to circumstances beyond the agency's control.

ADDRESSES: Copies are available from the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857, 301–443–1751.

FOR FURTHER INFORMATION CONTACT:

Donna M. Combs, Committee Management Office (HFA–306), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–443–2765.

SUPPLEMENTARY INFORMATION: Under section 13 of the Federal Advisory Committee Act (5 U.S.C. app. 2) and 21 CFR 14.60(c), FDA has filed with the Library of Congress the annual reports for the following FDA advisory committees that held closed meetings during the period October 1, 1993, through September 30, 1994:

Center for Biologics Evaluation and Research: Biological Response Modifiers Advisory Committee, Blood Products Advisory Committee, Vaccines and Related Biological Products Advisory Committee.

Center for Drug Evaluation and Research: Anesthetic and Life Support Drugs Advisory Committee, Anti-Infective Drugs Advisory Committee, Antiviral Drugs Advisory Committee, Cardiovascular and Renal Drugs Advisory Committee, Dermatologic and Ophthalmic Drugs Advisory Committee (formerly Dermatologic Drugs Advisory Committee), Gastrointestinal Drugs Advisory Committee, Nonprescription Drugs Advisory Committee, Oncologic Drugs Advisory Committee, Pulmonary-Allergy Drugs Advisory Committee.

Center for Devices and Radiological Health: Medical Devices Advisory Committee (consisting of reports for the Anesthesiology and Respiratory Therapy Devices Panel; Circulatory System Devices Panel; Clinical Chemistry and Clinical Toxicology Devices Panel (met jointly with the Microbiology Devices Panel); Dental Products Panel; Ear, Nose, and Throat Devices Panel; Gastroenterology and Urology Devices Panel; General and Plastic Surgery Devices Panel; General Hospital and Personal Use Devices Panel; Hematology and Pathology Devices Panel; Immunology Devices Panel; Neurological Devices Panel; Obstetrics and Gynecology Devices Panel; Ophthalmic Devices Panel; Orthopedic and Rehabilitation Devices Panel; and the Radiological Devices Panel).

Center for Veterinary Medicine: Veterinary Medicine Advisory Committee.

Office of Science: Science Board to the Food and Drug Administration.

National Center for Toxicological Research: Science Advisory Board to the National Center for Toxicological Research.

Annual reports are available for public inspection at: (1) The Library of Congress, Madison Bldg., Newspaper and Current Periodical Reading Room, 101 Independence Ave. SE., rm. 133, Washington, DC; and (2) the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857, between 9 a.m. and 4 p.m., Monday through Friday.

Dated: October 3, 1995.

David A. Kessler,

Commissioner of Food and Drugs.

[FR Doc. 95–25072 Filed 10–10–95; 8:45 am]

BILLING CODE 4160–01–F

National Institutes of Health

Government-Owned Inventions; Availability for Licensing

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

The inventions listed below are owned by agencies of the U.S. Government and are available for licensing in the U.S. in accordance with 35 U.S.C. 207 to achieve expeditious commercialization of results of federally funded research and development. Foreign patent applications are filed on selected inventions to extend market coverage for U.S. companies and may also be available for licensing.

ADDRESSES: Licensing information and copies of the U.S. patent applications listed below may be obtained by writing to the indicated Licensing Specialist at the Office of Technology Transfer,