August 11, 1995, the following corrections are made:

1. On page 41556, in footnote 89, ''1588'' is corrected to read ''1558,'' and on the same page, in footnote 90, in line 1, ''MDG'' is corrected to read ''MDB''.

2. On page 41557, in footnote 91, in line 1, the phrase ''of behavioral dependence'' is corrected to read ''of and behavioral dependence''.

3. On page 41558, in footnote 93, in line 4, ''*Parmacol. Biochem. Behav.*'' is corrected to read ''*Pharmacol. Biochemistry & Behav.*''

4. On page 41560, in footnote 101, in line 4, ''Page 50'' is corrected to read ''Pages 50–51.''

5. On page 41561, in footnote 105, in line 2, ''231–234'' is corrected to read ''231–241.''

6. On page 41588, in footnote 172, in line 8, ''12641–46'' is corrected to read ''02641–02646''.

7. On page 41621, in footnote 240a, in line 13, ''July 25, 2995'' is corrected to read ''July 25, 1995).''

Dated: December 12, 1995.

William B. Schultz,

Deputy Commissioner for Policy.

[FR Doc. 95–30745 Filed 12–18–95; 8:45 am]

BILLING CODE 4160–01–F

Public Health Service

Food and Drug Administration

Statement of Organization, Functions, and Delegations of Authority

Part H, Chapter HF (Food and Drug Administration) of the Statement of Organization, Functions, and Delegations of Authority for the Department of Health and Human Services (35 FR 3685, February 25, 1970, and 56 FR 29484, June 27, 1991, as amended most recently in pertinent part at 59 FR 17106, April 11, 1994) is amended to reflect the following reorganization within the Center for Devices and Radiological Health (CDRH), Office of Operations, Food and Drug Administration (FDA).

The Center for Devices and Radiological Health is abolishing the Office of Health Physics (OHP), the Office of Health Affairs (OHA), and the Office of Standards and Regulations (OSR) and realigning their functions into existing line and staff offices within the Center. The goal of this realignment is to more effectively manage the resources invested in these functional areas, consolidate similar functions, realign medical expertise closer to program needs, and streamline the current organizational structure.

Under section HF-B, Organization:

1. Delete subparagraphs Office of Health Physics (HFW12), the Office of Health Affairs (HFW13), and the Office of Standards and Regulations (HFW14) under paragraph Center for Devices and Radiological Health (HFW), in their entirety.

2. Insert the following new subparagraphs under paragraph *Office of Operations (HFA9), Center for Devices and Radiological Health (HFW)* reading as follows:

Office of Systems and Management (HFW11). Advises the Center Director regarding all administrative management matters.

Plans, develops, and implements Center management policies and programs concerning financial and human resource management, contracts and grants management, conference management, occupational safety, organizational, and general office services support.

Develops and implements the Center's long-range, strategic, and operational plans.

Develops and applies evaluation techniques to measure the effectiveness of Center programs.

Provides general information and technical publication services to the Center.

Plans, conducts, and coordinates Center committee management activities.

Determines and implements Center strategy and utilization of information management resources.

Designs administrative, scientific, and technical information systems in support of Center programs.

Provides assistance to Center staff in accessing information necessary to carry out the Center's mission.

Coordinates requests and Center activities pertaining to the Freedom of Information and Privacy Acts.

Office of Health and Industry Programs (HFWG). Analyzes medical device and radiation-emitting product user-related problems and conducts research, applying systems analysis and human factors to problem identification and solution strategies. Implements and evaluates user-related solution strategies.

Conducts and evaluates programs to provide technical and other nonfinancial assistance to small manufacturers of medical devices to promote their understanding of compliance with the medical device amendments and regulations.

Provides, maintains, and applies expertise in communications technology in support of Center and FDA programs.

Develops and implements strategies for obtaining, analyzing, and incorporating the views and needs of health professionals, lay device users, and industry into the Center policy and decision-making processes as well as in problem analysis, resolution strategy development, implementation, and evaluation processes.

Establishes and operates a program to implement the Mammography Quality Standards Act of 1992.

Provides leadership and technical expertise to the Center and other Departmental components in applying health physics procedures and radiation protection principles.

Advises the Center Director and appropriate Agency officials on FDA regulation development responsibilities relating to medical devices and radiological health activities. Serves as the Center focal point for liaison on regulations development activities with the Office of General Counsel.

Coordinates the development, review and submission of Federal Register publications for the Center. Prepares position statements for the Center on standards promulgated by other organizations.

Coordinates international relations activities as required by the Safe Medical Devices Act of 1990.

Office of Science and Technology (HFWE). Provides scientific support and laboratory analyses in response to the program needs of other Center and Agency components.

Plans, develops, and implements an intramural science program covering key areas of engineering, physics, and biology; develops, modifies, and validates test methods and measurement techniques, risk assessments and hazard analyses, and generic techniques to enhance product safety and usefulness.

Provides scientific and engineering support in the review of regulatory documents, the development of regulatory decisions, and the analysis of postmarket surveillance issues.

Plans, conducts, or stimulates research on the human health effects of radiation and medical devices.

Participates in the development of national and international consensus standards and voluntary guidelines through interaction with appropriate standards committees; coordinates with other standards-setting groups representing national and international standards-setting organizations; conducts the review and analysis of performance standards, guides and documents related to the Center's mission.

Establishes official liaisons with Standards Development Organizations. Coordinates the liaison within the Center. Establishes and maintains

records on committee participation and status of medical device standards used by the Center.

3. Prior Delegations of Authority. Pending further delegations, directives, or orders by the Commissioner of Food and Drugs, all delegations of authority to positions of the affected organizations in effect prior to this date shall continue in effect in them or their successors.

Dated: November 28, 1995.

David A. Kessler,

Commissioner of Food and Drugs.

[FR Doc. 95–30813 Filed 12–18–95; 8:45 am]

BILLING CODE 4160–01–F

Food and Drug Administration

Statement of Organization, Functions, and Delegations of Authority

Part H, Chapter HF (Food and Drug Administration) of the Statement of Organization, Functions, and Delegations of Authority for the Department of Health and Human Services (35 FR 3685, February 25, 1970, and 56 FR 29484, June 27, 1991, as amended most recently in pertinent part at 60 FR 53382, October 13, 1995) is amended to reflect the change in title of the International Affairs Staff to the Office of International Affairs, Office of External Affairs (OEA), in the Food and Drug Administration (FDA).

The Office of International Affairs will continue to serve as the Agency focal point for developing and maintaining international communications and programs. FDA believes that the increase in international activity with regard to FDA regulated products and activities necessitates the elevation of the International Affairs Staff to office level status within OEA and that this action further enhances the management and coordination of Agency international activities.

Under section HF-B, Organization:

1. Delete the subparagraph, *International Affairs Staff (HFAQA),* under the *Office of External Affairs (HFAQ),* in its entirety and insert a new subparagraph, Office of International Affairs (HFAQA), under the Office of External Affairs (HFAQ), reading as follows:

Office of International Affairs (HFAQA). Serves as the Agency focal point for developing and maintaining international communications and programs.

Establishes and provides an Agency liaison on international activities with the Department, Public Health Service (PHS), and other Federal agencies, foreign governments, including foreign embassies, and international organizations.

Represents the Agency at meetings, conferences, and symposia relating to international obligations; briefs Agency participants in such international activities.

Establishes, identifies, interprets, and clarifies, in cooperation with appropriate Agency components, the Agency's international obligations and needs, including those associated with bilateral programs which involve extra budgetary support.

Establishes and maintains an international information exchange program concerning Agency policies and programs to provide interchange between FDA and counterpart agencies in foreign countries and international organizations.

Assists in the development, negotiation, and monitoring of agreements with foreign governments and international organizations in cooperation with appropriate Agency components; and acts as the Agency focal point for intergovernmental conferences.

Negotiates the preparation and implementation of technical assistance programs (including formal training programs and surveys) with foreign governments and international organizations in areas relating to the Agency mission. Coordinates ongoing technical assistance operations with appropriate components within the Department, PHS, and the Agency.

Directs the Agency's International Visitors Program, providing participants with policy briefings, technical training, and/or assistance in response to specific needs

2. Prior Delegations of Authority. Pending further delegations, directives, or orders by the Commissioner of Food and Drugs, all delegations of authority to positions of the affected organizations in effect prior to this date shall continue in effect in them or their successors.

Dated: December 5, 1995.

David A. Kessler,

Commissioner of Food and Drugs.

[FR Doc. 95–30744 Filed 12–18–95; 8:45 am]

BILLING CODE 4160–01–F

Health Resources and Services Administration

Advisory Council; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), announcement is made of the following National Advisory body scheduled to meet during the month of January 1996.

Name: Council on Graduate Medical Education

Date and Time: January 9, 1996, 1:00 p.m.–5:00 p.m., January 10, 1996, 8:30 a.m.–4:00 p.m.

Place: Governor's House Hotel, 17th Street at Rhode Island Avenue, N.W., Washington, D.C. 20036

The meeting is open to the public.

Purpose: Provides advice and recommendations to the Secretary and to the Committees on Labor and Human Resources, and Finance of the Senate and the Committees on Energy and Commerce and Ways and Means of the House of Representatives, with respect to (A) the supply and distribution of physicians in the United States; (B) current and future shortages of physicians in medical and surgical specialties and subspecialties; (C) issues relating to international medical graduates; (D) appropriate Federal policies regarding (A), (B), and (C) above; (E) appropriate efforts to be carried out by medical and osteopathic schools, public and private hospitals and accrediting bodies regarding matters in (A), (B), and (C) above; (F) deficiencies in the needs for improvements in, existing data bases concerning supply and distribution of, and training programs for physicians in the United States.

Agenda: The Agenda will include a panel to discuss International Medical Graduates, entry and participation in the U.S. physician workforce. A panel to discuss Legislation and GME Reform, House Ways and Means Committee. Report on transition funding issues; Report and updates on the work groups, Minorities in Medicine; Geographic Distribution/Medical Education Consortia; Physician Competencies in a Managed Care World; and IMG Entry and Participation in the Physician Workforce.

Anyone requiring information regarding the meeting should contact F. Lawrence Clare, M.D., M.P.H., Deputy Executive Secretary, Telephone 301–443–6326, Council on Graduate Medical Education, Division of Medicine, Bureau of Health Professions, Room 9A–27, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857.

Agenda items are subject to change as priorities dictate.

Dated: December 12, 1996.

Jackie E. Baum,

Advisory Committee Management Officer.

[FR Doc. 95–30816 Filed 12–18–95; 8:45 am]

BILLING CODE 4160–15–P

National Institutes of Health

National Institute of Mental Health; Notice of Closed Meeting

Pursuant to Section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of the following meeting