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 AnnMarie Williams, Conference Management Staff, at 301–594–1283, ext. 113, 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: June 26, 2002.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning, and Legislation.

[FR Doc. 02–16904 Filed 7–5–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Obstetrics and Gynecology Devices Panel of the Medical Devices 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). At least one portion of the meeting will be closed to the public.

Name of the Committee: Obstetrics and Gynecology Devices Panel of the Medical Devices 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 July 22, 2002, from 8:30 a.m. to 5 p.m., and on July 23, 2002, from 8 a.m. to 11 a.m.

Location: DoubleTree Hotel, Plaza I and II, 1750 Rockville Pike, Rockville, MD.

Contact Person: Joyce M. Whang, Center for Devices and Radiological Health (HFZ–470), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–594–1180, or FDA Advisory Committee Information Hotline, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12524. Please call the Information Line for up-to-date information on this meeting.

Agenda: On July 22, 2002, the committee will hear a presentation on post-market surveillance of vacuum assisted delivery devices. The committee will also discuss, make recommendations, and vote on a premarket approval application for a permanent contraceptive device. Background information, including the agenda and questions for the committee, will be available to the public 1 business day before the meeting on the Internet at http://www.fda.gov/cdrh/panelmtg.html. Material for the July 22, 2002, session will be posted on July 19, 2002.

Procedure: On July 22, 2002, from 8:30 a.m. to 5 p.m., the meeting is open to the public. 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 July 11, 2002. Oral presentations from the public will be scheduled between approximately 9:30 a.m. and 10 a.m., and 3:30 p.m. and 4 p.m. on July 22, 2002. Time allotted for each presentation may be limited. Those desiring to make formal presentations should notify the contact person before July 11, 2002, 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.

Closed Committee Deliberations: On July 23, 2002, from 8 a.m. to 11 a.m., the meeting will be closed to permit discussion and review of trade secret and/or confidential commercial information (5 U.S.C. 552b(c)(4)) regarding pending and future device issues. In addition, the committee will discuss and review trade secret and/or confidential commercial information presented by a sponsor.

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 AnnMarie Williams, Conference Management Staff, at 301–594–1283, ext. 113, at least 7 days in advance of the meeting.

FDA regrets that it was unable to publish this notice 15 days prior to the July 22, 2002, Obstetrics and Gynecology Devices Panel of the Medical Devices Advisory Committee meeting. Because the agency believes there is some urgency to bring these issue(s) to public discussion and qualified members of the Obstetrics and Gynecology Devices Panel of the Medical Devices Advisory Committee were available at this time, the Commissioner of Food and Drugs concluded that it was in the public interest to hold this meeting even if there was not sufficient time for the customary 15-day public notice.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: July 2, 2002.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning, and Legislation.

[FR Doc. 02–17115 Filed 7–5–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02D–0274]

Denture Cleaners, Adhesives, Cushions, and Repair Materials; Revocation of Compliance Policy Guide 7124.05

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is revoking the Compliance Policy Guide (CPG) entitled ''Sec. 315.200 Status of Dental Supplies such as Denture Cleaners, Adhesives, Cushions, and Repair Materials as a Device or Cosmetic (CPG 7124.05).'' This CPG is no longer necessary because the agency has classified these products as devices.

DATES: The revocation is effective August 7, 2002.

ADDRESSES: Submit written requests for single copies of the CPG to the Division of Compliance Policy (HFC–230), Office of Enforcement, Office of Regulatory Affairs, Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857 (301–827–0411) or fax your request to 301–827–0482.

A copy of the CPG may be seen in the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, between 9 a.m. and 4 p.m., Monday through Friday.

FOR FURTHER INFORMATION CONTACT: Jeffrey B. Governale, Division of Compliance Policy (HFC–230), Office of Enforcement, Office of Regulatory Affairs, Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–0411.

SUPPLEMENTARY INFORMATION:

I. Background

FDA issued the CPG entitled ''Sec. 315.200 Status of Dental Supplies such

as Denture Cleaners, Adhesives, Cushions, and Repair Materials as a Device or Cosmetic (CPG 7124.05)'' on April 26, 1976. This CPG, as revised on August 9, 1988, considered these products to be devices within the meaning of section 201(h) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 321(h)).

In accordance with section 513 of the act (21 U.S.C. 360c), the agency has classified dental products as devices by regulation, including but not limited to:

1. Karaya and sodium borate with or without acacia denture adhesive (21 CFR 872.3400)
2. Ethylene oxide homopolymer and/or carboxymethylcellulose sodium denture adhesive (21 CFR 872.3410)
3. Carboxymethylcellulose sodium and cationic polyacrylamide polymer denture adhesive (21 CFR 872.3420)
4. Ethylene oxide homopolymer and/or karaya denture adhesive (21 CFR 872.3450)
5. Polyacrylamide polymer (modified cationic) denture adhesive (21 CFR 872.3480)
6. Carboxymethylcellulose sodium and/or polyvinylmethylether maleic acid calcium-sodium double salt denture adhesive (21 CFR 872.3490)
7. Polyvinylmethylether maleic anhydride (PVM–MA), acid copolymer, and carboxymethylcellulose sodium (NACMC) denture adhesive (21 CFR 872.3500)
8. Over-the-counter (OTC) denture cleanser (21 CFR 872.3520)
9. Mechanical denture cleaner (21 CFR 872.3530)
10. OTC denture cushion or pad (21 CFR 872.3540)
11. OTC denture repair kit (21 CFR 872.3570)
12. Denture relining, repairing, or rebasing resin (21 CFR 872.3760)

Given these device classifications, FDA is revoking CPG 7124.05, in its entirety, to eliminate unnecessary compliance policy.

II. Electronic Access

Prior to August 7, 2002, a copy of the CPG may also be downloaded to a personal computer with access to the Internet. The Office of Regulatory Affairs (ORA) home page includes the referenced document that may be accessed at http://www.fda.gov/ora/compliance_ref/cpg/cpgdev/cpg315–200.html.

Dated: June 28, 2002.

Dennis E. Baker,

Associate Commissioner for Regulatory Affairs.

[FR Doc. 02–17079 Filed 7–5–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Council on Graduate Medical Education; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Public Law 92–463), announcement is made of the following National Advisory body scheduled to meet during the month of September:

Name: Council on Graduate Medical Education (COGME).

Date and Time: September 11, 2002, 8:30 a.m.–4:30 p.m., September 12, 2002, 8 a.m.–11:15 a.m.

Place: Holiday Inn Select, Versailles 1, 8120 Wisconsin Avenue, Bethesda, MD 20814.

The meeting is open to the public.

Agenda: The agenda for September 11 will include: Welcome and opening comments from the Associate Administrator for Health Professions, the Chair of COGME, and the Acting Executive Secretary of COGME. There will be a panel of speakers on the topic of ''Competencies in Graduate Medical Education'' and a panel of speakers on the topic of ''Financial Situation of Teaching Hospitals.''

In the afternoon the Council's three workgroups will convene. They are: Workgroup on Diversity, Workgroup on Graduate Medical Education Financing, and Workgroup on Workforce.

The agenda for September 12 will include a discussion of the June 17–18 Health Professions Education Summit co-sponsored by the Council on Graduate Medical Education (COGME), the National Advisory Council on Nurse Education and Practice, and the Institute of Medicine. The three workgroup chairs will give their reports. There will be a discussion of the status of COGME's 2002 Summary Report, plans for future work, and new business.

Anyone requiring information regarding the meeting should contact Stanford M. Bastacky, D.M.D., M.H.S.A., Acting Executive Secretary, Council on Graduate Medical Education, Division of Medicine and Dentistry, Bureau of Health Professions, Room 9A–27, Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857, Telephone (301) 443–6326.

Agenda items are subject to change as priorities dictate.

Dated: July 2, 2002.

Jane M. Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 02–17047 Filed 7–5–02; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Committee on Training in Primary Care Medicine and Dentistry; Notice of Meeting

In accordance with section 10 (a) (2) of the Federal Advisory Committee Act (Public Law 92–463), announcement is made of the following National Advisory body scheduled to meet during the month of August 2002.

Name: Advisory Committee on Training in Primary Care Medicine and Dentistry (ACTPCMD).

Date and Time: August 19, 2002; 8:30 a.m.–4:45 p.m.

Place: The Holiday Inn, 8120 Wisconsin Avenue, Bethesda, Maryland 20814.

The meeting is open to the public.

Purpose: The Advisory Committee on Training in Primary Care Medicine and Dentistry (ACTPCMD) will discuss and consider criteria for performance measurement and outcome assessment of primary care medicine and dentistry grant programs that are funded through Title VII, section 747 of the Public Health Service Act, as amended. The Committee will review current performance measurement and outcome assessment methods, discuss potential alternative methods, and consider recommendations of potential performance measurement and outcome assessment methods that might be employed in the future.

Agenda: The meeting on Monday, August 19 will begin with welcoming and opening comments from the Chair and Executive Secretary. A plenary session will follow in which Division of Medicine and Dentistry staff will review criteria currently used to measure performance and assess the outcome of primary care medicine and dentistry grant programs funded through Title VII, section 747 of the Public Health Service Act, as amended. Following this presentation, Committee members will discuss the criteria currently used, consider potential alternative criteria for performance measurement and outcome assessment of the aforementioned grant programs, and formulate recommendations for alternative criteria.

Anyone interested in obtaining a roster of members or other relevant information should write or contact Stan Bastacky, D.M.D., M.H.S.A., Deputy Executive Secretary, Advisory Committee on Training in Primary Care Medicine and Dentistry, Parklawn Building, Room 9A–21, 5600 Fishers Lane, Rockville, Maryland 20857, Telephone (301) 443–6326. The Web address for information on the Advisory Committee is *http://www.bhpr.hrsa.gov/dm/actpcmd.htm.*