DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Orthopaedic and Rehabilitation 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). The meeting will be open to the public.

Name of Committee: Orthopaedic and Rehabilitation 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 November 20 and 21, 2002, from 9:30 a.m. to 5 p.m.

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

Contact Person: Hany W. Demian, Center for Devices and Radiological Health (HFZ–410), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–594–2036, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12521. Please call the Information Line for up-to-date information on this meeting.

Agenda: On November 20, 2002, the committee will discuss, make recommendations, and vote on a premarket approval application (PMA) for a stair-climbing wheelchair. On November 21, 2002, the committee will discuss, make recommendations, and vote on a PMA for growth factors soaked in a collagen sponge used to treat tibial fractures. In addition, on November 21, 2002, the committee will have a general discussion on preclinical and clinical data for spinal devices. Background information for each day's topic, 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 November 20, 2002, session will be posted on November 19, 2002; material for the November 21, 2002, session will be posted on November 20, 2002.

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 November 15, 2002. On each day, oral presentations from the public will be scheduled for approximately 30 minutes at the beginning of each PMA topic and for approximately 30 minutes near the end of the committee deliberations. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before November 15, 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.

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 November 20 and 21, 2002, Orthopaedic and Rehabilitation Devices Panel of the Medical Devices Advisory Committee meeting. Because the agency believes there is some urgency to bring these issues to public discussion and qualified members of the Orthopaedic and Rehabilitation 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: November 5, 2002.

Linda Arev Slkadany,

 $Senior\ Associate\ Commissioner\ for\ External\ Relations.$

[FR Doc. 02–28853 Filed 11–12–02; 8:45 am] BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Pulmonary-Allergy Drugs 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: Pulmonary-Allergy Drugs 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 December 20, 2002, from 7:30 a.m. to 5 p.m.

Location: Holiday Inn, The Ballrooms, Two Montgomery Village Ave.,

Gaithersburg, MD.

Contact Person: Kimberly Littleton Topper, Center for Drug Evaluation and Research (HFD–21), Food and Drug Administration, 5600 Fishers Lane (for express delivery, 5630 Fishers Lane, rm. 1093), Rockville, MD 20857, 301–827–7001, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12545. Please call the Information Line for up-to-date information on this meeting.

Agenda: The committee will discuss new drug application (NDA) 20–959, Ebastine by Almirall Prodesfarma, for the proposed indication of relief of nasal and nonnasal symptoms associated with seasonal and perennial allergic rhinitis in adults and children 12 years of age and older.

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 December 13, 2002. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before December 13, 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.

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 Kimberly Topper 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: November 4, 2002.

Linda Arey Skladany,

Senior Associate Commissioner for External Relations.

[FR Doc. 02–28852 Filed 11–12–02; 8:45 am]

BILLING CODE 4160-01-S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget, in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to OMB for review, call the HRSA Reports Clearance Office on (301) 443–1129.

The following request has been submitted to the Office of Management and Budget for review under the Paperwork Reduction Act of 1995:

Proposed Project: The Health Professions Student Loan (HPSL) and Nursing Student Loan (NSL) Programs: Forms—(OMB No. 0915–0044)— Revision

The HPSL Program Provides longterm, low-interest loans to students

attending schools of medicine, osteopathic medicine, dentistry, veterinary medicine, optometry, podiatric medicine, and pharmacy. The NSL Program provides long-term, lowinterest loans to students who attend eligible schools of nursing in programs leading to a diploma in nursing, and an associate degree, a baccalaureate degree, or a graduate degree in nursing. Participating HPSL and NSL schools are responsible for determining eligibility of applicants, making loan, and collecting monies owed by borrowers on their outstanding loans. The deferment form (HRSA form 519) provides the schools with documentation of a borrower's eligibility for deferment. The Annual Operating Report (AOR-HRSA form 501) provides the Federal Government with information from participating and non-participating schools (schools that are no longer granting loans but are required to report and maintain program records, student records, and repayment records until all student loans are repaid in full and all monies due the Federal Government are returned) relating to HPSL and NSL program operations and financial activities.

The estimate of burden for the forms are as follows:

Form and number	Number of respondents	Responses per respondent	Total responses	Hours per responses	Total burden hours
Defer-HRSA-519AOR-HRSA-501	6,000 1,048	1 1	6,000 1,048	10 min 4 hrs.	1,000 4,192
Total Burden	7,048		7,048		5,192

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to: John Morrall, Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: November 6, 2002.

Jane M. Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 02-28855 Filed 11-12-02; 8:45 am]

BILLING CODE 4165-15-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Children's Hospitals Graduate Medical Education (CHGME) Payment Program

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Notice of children's hospitals graduate medical education (CHGME) Payment Program conference calls.

SUMMARY: This document announces scheduled CHGME Payment Program conference calls for calendar year 2003. The purpose of these conference calls is to provide technical assistance related to the CHGME Payment Program.

DATES: The conference calls will be held on Wednesday, January 22, 2003, from 1:30 p.m. to 3:30 p.m. e.s.t., Wednesday, April 23, 2003, from 1:30 p.m. to 3:30 p.m. e.s.t., and Wednesday, October 22, 2003, from 1:30 p.m. to 3:30 p.m. e.s.t.

FOR FURTHER INFORMATION CONTACT:

Ayah E. Johnson, Ph.D., telephone: (301) 443–1058; Division of Medicine and Dentistry, Bureau of Health Professions, Health Resources and Services Administration, Parklawn Building, 5600 Fishers Lane, Room 9A–27, Rockville, Maryland 20857; or by e-mail at: ajohnson@hrsa.gov.

SUPPLEMENTARY INFORMATION: The CHGME Payment Program, as authorized by section 340E of the Public Health Service (PHS) Act (the Act) (42 U.S.C. 256e), provides funds to children's hospitals to address disparity in the level of Federal funding for children's hospitals that result from Medicare funding for graduate medical education (GME). Pub. L. 106–310 amended the CHGME statute to extend the program through fiscal year (FY) 2005.

The statute authorized \$280 million for both direct and indirect medical education payments in FY 2000, \$285 million in FY 2001, and for each of the