

early childhood community, formulate the OHS strategic plans and long-term goals, provide guidance and support with budget planning, oversee development of regulations and other policy issuances, and serve as liaison with ACF and HHS legislative offices on all Congressional matters relating to Head Start.

F. Grants and Contracts Division (KUG): The Grants and Contracts Division (GCD) (1) Oversees matters related to competitive funding opportunities; (2) manages competition, paneling, and selection of national contracts and Head Start and Early Head Start replacement grantees; (3) provides ongoing fiscal oversight of national contracts; (4) serves as the lead for the OHS Program Management and Fiscal Operations Center; and (5) serves as the liaison to the Office of Administration, Divisions of Grants Management and Division of Grants Policy.

G. State Initiatives Division (KUH): The State Initiatives Division (SID) leads and consolidates collaboration efforts to new and expanding Head Start programs. The Division will promote collaborations with state pre-k programs, local child care providers and other national and state early childhood efforts to ensure the sustainability of strong collaborations. The Division serves as the locus for ensuring that mandates in the Head Start Act regarding collaboration are implemented as well as coordination with the U.S. Department of Education and state early childhood entities. The Division will focus on State Advisory Councils, Centers of Excellence, State Collaboration Offices, and the Training and Technical Assistance System.

II. *Continuation of Policy.* Except as inconsistent with this reorganization, all statements of policy and interpretations with respect to organizational components affected by this notice within ACF, heretofore issued and in effect on this date of this reorganization are continued in full force and effect.

III. *Delegation of Authority.* All delegations and redelegations of authority made to officials and employees of affected organizational components will continue in them or their successors pending further redelegations, provided they are consistent with this reorganization.

IV. *Funds, Personnel, and Equipment.* Transfer of organizations and functions affected by this reorganization shall be accompanied in each instance by direct and support funds, positions, personnel, records, equipment, supplies, and other resources.

This reorganization will be effective upon date of signature.

Dated: December 16, 2010.

David A. Hansell,

Acting Assistant Secretary for Children and Families.

[FR Doc. 2010-32462 Filed 12-23-10; 8:45 am]

BILLING CODE 4184-40-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2010-N-0001]

Circulatory System 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: Circulatory System 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 January 25 and 26, 2011, from 8 a.m. to 6 p.m.

Location: Holiday Inn, Main Ballroom, Two Montgomery Village Ave., Gaithersburg, MD. Information regarding special accommodations due to a disability, visitor parking and transportation may be accessed at: <http://www.fda.gov/AdvisoryCommittees/default.htm>; under the heading "Resources for You," click on "White Oak Conference Center Parking and Transportation Information for FDA Advisory Committee Meetings." Please note that visitors to the White Oak Campus must enter through Building 1.

Contact Person: James Swink, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Silver Spring, MD 20993, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 3014512625. Please call the Information Line for up-to-date information on this meeting. A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's Web site and call the appropriate advisory committee hot

line/phone line to learn about possible modifications before coming to the meeting.

Agenda: On January 25, 2011, the committee will discuss and make recommendations regarding regulatory classification of Automated External Defibrillators to either reconfirm to class III (subject to premarket approval application (PMA)) or reclassify to class II (subject to premarket notification (510(k))), as directed by section 515(i) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C 360e(i)).

On January 26, 2011, the committee will discuss, make recommendations and vote on information related to the PMA supplement for the RX Acculink Carotid Stent System, sponsored by Abbott Vascular. The RX Acculink is indicated for treatment of patients at high and standard risk for adverse events from carotid endarterectomy who require carotid revascularization and meet the criteria outlined as follows:

1. Patients with neurological symptoms and >50 percent stenosis of the common or internal carotid artery or patients without neurological symptoms and >80 percent (high risk) or >70 percent (standard risk) stenosis of the common or internal carotid artery and

2. Patients must have a reference vessel diameter within the range of 4.0 and 9.0 mm at the target lesion.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at <http://www.fda.gov/AdvisoryCommittees/Calendar/default.htm>. Scroll down to the appropriate advisory committee link.

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 on or before January 18, 2011. Oral presentations from the public will be scheduled for 1 hour at approximately 1 p.m., immediately following lunch on both days. Those individuals interested in making formal oral presentations should notify the contact person 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 on

or before January 10, 2011. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by January 11, 2011.

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, 301-796-5966, at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at <http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm> for procedures on public conduct during advisory committee meetings.

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

Dated: December 20, 2010.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2010-32367 Filed 12-23-10; 8:45 am]

BILLING CODE 4160-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2010-N-0001]

Oncologic Drugs Advisory Committee; Cancellation

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The meeting of the Oncologic Drugs Advisory Committee scheduled for February 9, 2011, is cancelled. This meeting was announced in the **Federal Register** of December 6, 2010 (75 FR 75680). On February 9, 2011, the Oncologic Drugs Advisory Committee was scheduled to discuss biologics license application (BLA) 125377, with the proposed trade name YERVOY (ipilimumab), submitted by Bristol-

Myers Squibb Co. The proposed indication (use) for this product is for the treatment of advanced melanoma in patients who have received prior therapy. This meeting has been cancelled because the issues for which FDA was seeking the scientific input of the committee have been resolved.

FOR FURTHER INFORMATION CONTACT:

Nicole Vesely, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, rm. 2417, Silver Spring, MD 20993-0002, 301-796-9001, FAX: 301-847-8533, e-mail: Nicole.vesely@fda.hhs.gov, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), and follow the prompts to the desired center or product area. Please call the Information Line for up-to-date information on this meeting.

Dated: December 21, 2010.

Jill Hartzler Warner,

Acting Associate Commissioner for Special Medical Programs.

[FR Doc. 2010-32413 Filed 12-23-10; 8:45 am]

BILLING CODE 4160-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Eunice Kennedy Shriver National Institute of Child Health & Human Development; Notice of Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of a meeting of the National Advisory Child Health and Human Development Council.

The meeting will be open to the public, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and/or contract proposals and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications and/or contract proposals, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Advisory Child Health and Human Development Council.

Date: January 20, 2011.

Open: 8 a.m. to 12:20 p.m.

Agenda: (1) A report of the Director, NICHD; (2) Select Pay Plan Presentation; (3) NIH Peer Review Enhancement Update, (4) NIH Support for Human Embryonic Stem Cell Research Update.

Place: National Institutes of Health, Building 31, 31 Center Drive, C-Wing, Conference Room 6, Bethesda, MD 20892.

Closed: 12:20 p.m. to 5 p.m.

Agenda: To review and evaluate grant applications and/or proposals.

Place: National Institutes of Health, Building 31, 31 Center Drive, C-Wing, Conference Room 6, Bethesda, MD 20892.

Contact Person: Yvonne T. Maddox, PhD, Deputy Director, National Institute of Child Health and Human Development, NIH, 9000 Rockville Pike MSC 7510, Building 31, Room 2A03, Bethesda, MD 20892. (301) 496-1848.

Any interested person may file written comments with the committee by forwarding the statement to the Contact Person listed on this notice. The statement should include the name, address, telephone number and when applicable, the business or professional affiliation of the interested person.

In the interest of security, NIH has instituted stringent procedures for entrance onto the NIH campus. All visitor vehicles, including taxis, hotel, and airport shuttles will be inspected before being allowed on campus. Visitors will be asked to show one form of identification (for example, a government-issued photo ID, driver's license, or passport) and to state the purpose of their visit.

Information is also available on the Institute's/Center's home page: <http://www.nichd.nih.gov/about/nachhd.htm>, where an agenda and any additional information for the meeting will be posted when available.

In order to facilitate public attendance at the open session of Council, reserve seating will be made available to the first five individuals reserving seats in the main meeting room, Conference Room 6. Please contact Ms. Lisa Kaeser, Program and Public Liaison Office, NICHD, at 301-496-0536 to make your reservation. Additional seating will be available in the meeting overflow rooms, Conference Rooms 7 and 8. Individuals will also be able to view the meeting via NIH Videocast. Please go to the following link for Videocast access instructions at: <http://www.author.nichd.nih.gov/about/overview/advisory/nachhd/virtual-meeting-201005.cfm>. The meeting is partially closed to the public.

(Catalogue of Federal Domestic Assistance Program Nos. 93.864, Population Research; 93.865, Research for Mothers and Children; 93.929, Center for Medical Rehabilitation Research; 93.209, Contraception and Infertility Loan Repayment Program, National Institutes of Health, HHS)