DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): NIOSH Occupational Safety and Health Project Grants, Program Announcement Number (PAR) 06–484

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the aforementioned meeting:

Time and Date: 8:30 a.m.–5:30 p.m., March 15, 2007 (Closed).

Place: Residence Inn, 1456 Duke Street, Alexandria, VA 22314, telephone 703–548–5474.

Status: The meeting will be closed to the public in accordance with provisions set forth in section 552b(c)(4) and (6), Title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92–463.

Matters To Be Discussed: The SEP meeting will include the review, discussion, and evaluation of applications received in response to ''NIOSH Occupational Safety and Health Project Grants,'' PAR 06–484. The applications being reviewed include information of a confidential nature, including personal information concerning individuals associated with the applications.

This **Federal Register** Notice is being published on less than 15 calendar days notice to the public (41 CFR 102–3.150(b)), for the following reason: The cancellation of a preparatory meeting scheduled for January 16th due to inclement weather caused the late publication of this notice. Convening the preparatory meeting was necessary before this meeting could be scheduled. The preparatory meeting occurred on February 20–21, 2007, which enabled the program to finalize plans for this meeting.

For Further Information Contact: Charles Rafferty, Ph.D., Designated Federal Officer, 1600 Clifton Road NE, Atlanta, GA 30333, telephone 404–498–2582.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both CDC and the Agency for Toxic Substances and Disease Registry.

Dated: February 26, 2007.

Elaine L. Baker,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

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BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

The Essentials of Food and Drug Administration Medical Device Regulations: A Primer for Manufacturers and Suppliers; Public Seminar

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public seminar.

SUMMARY: The Food and Drug Administration's (FDA's) Center for Devices and Radiological Health and Office of Regulatory Affairs, in cooperation with AdvaMed's Medical Technology Learning Institute, is announcing a series of three seminars on FDA medical device regulations.

These 2-day seminars, which are designed to address the training needs of start up and small device manufacturers and their suppliers, will include both industry and FDA perspectives and a question and answer period.

Dates: The seminars are planned for the following dates:

1. March 15 and 16, 2007, in Irvine, CA 92614. Details about dates are posted on AdvaMed's Web site at: *www.advamed.org/irvine*.[1]

2. May 22 and 23, 2007, in Lakewood, CO 80228. Details about dates are posted on AdvaMed's Web site at: *www.advamed.org/denver*.

3. June 6 and 7, 2007, in Pittsburgh, PA, Details about dates are posted on AdvaMed's Web site at: *www.advamed.org/pittsburgh*.

Locations: The seminars are planned for the following locations:

1. March 15 and 16, 2007, Crown Plaza Hotel, 17941 Von Karman, Irvine, CA 92614. Details about location sites are posted on AdvaMed's Web site at: *www.advamed.org/irvine*.

2. May 22 and 23, 2007, Sheraton Denver West, 360 Union Blvd., Lakewood, CO 80228. Details about location sites are posted on AdvaMed's Web site at: *www.advamed.org/denver*.

3. June 6 and 7, 2007, Hilton Pittsburgh, 600 Commonwealth Pl., Pittsburgh, PA 15222, *www.HiltonPittsburgh.com*. Details about location sites are posted on AdvaMed's Web site at: *www.advamed.org/pittsburgh*.

Contact: *For FDA*: William Sutton, Division of Small Manufacturers, International and Consumer Assistance, Center for Devices and Radiological Health (HFZ–220), 1350 Piccard Dr., Rockville, MD 20850, 800–638–2041, ext. 125, FAX: 240–276–3151, e-mail: *William.sutton@fda.hhs.gov*.

For AdvaMed: Dia Black, 202–434–7231, FAX: 202–783–8750, e-mail: *DBlack@AdvaMed.org*.

Registration: The registration fee for FDA employees is waived. Send registration information (including name, title, firm name, address, telephone, and fax number) and the registration fee of $495 per person to AdvaMed contact Dia Black, 202–434–7231, FAX: 202–783–8750. Payment forms accepted are major credit card (MasterCard, Visa, or American Express) or company check. If you wish to pay by check, contact Dia Black at: *DBlack@AdvaMed.org*.

To register via the Internet, go to *www.AdvaMed.org*. The latest information on dates/venue sites will be posted on this Web site at: *www.advamed.org/irvine*, *www.advamed.org/denver*, and *www.advamed.org/pittsburgh* (FDA has verified the Web site addresses, but is not responsible for changes to the Web sites after this document publishes in the **Federal Register**).

For more information on the meeting, or for questions on registration, contact Dia Black (see *Contact*).

Attendees are responsible for their own accommodations. For further hotel information and driving directions, go to the registration Web site.

The registration fee will be used to offset the expenses of hosting the conference, including meals (breakfasts and a lunch), refreshments, meeting rooms, and training materials. It also includes a networking reception on the evening of the first day of each seminar.

Space is limited; therefore, interested parties are encouraged to register early. There will be no onsite registration.

If you need special accommodations due to a disability, please contact Dia Black (see *Contact*) at AdvaMed at least 7 days in advance of the seminar.

SUPPLEMENTARY INFORMATION: The ''Essentials of FDA Medical Device Regulations: A Primer for Manufacturers and Suppliers'' seminar helps fulfill the Department of Health and Human Services' and FDA's important mission to protect the public health by educating new entrepreneurs on the essentials of FDA device regulations. FDA has made education of the medical device community a high priority to assure the quality of products reaching the marketplace and to increase the rate of voluntary industry compliance with regulations.

[1] FDA has verified the Web site addresses, but FDA is not responsible for any subsequent changes to the Web sites after this document publishes in the **Federal Register**.