

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 15 people. Call the ATSDR/ORR field office for the conference bridge line and access code.

Background: A Memorandum of Understanding (MOU) was signed in October 1990 and renewed in September 2000 between ATSDR and DOE. The MOU delineates the responsibilities and procedures for ATSDR's public health activities at DOE sites required under sections 104, 105, 107, and 120 of the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA or ASuperfund®). These activities include health consultations and public health assessments at DOE sites listed on, or proposed for, the Superfund National Priorities List and at sites that are the subject of petitions from the public; and other health-related activities such as epidemiologic studies, health surveillance, exposure and disease registries, health education, substance-specific applied research, emergency response, and preparation of toxicological profiles.

In addition, under an MOU signed in December 1990 with DOE and replaced by an MOU signed in 2000, the Department of Health and Human Services (HHS) has been given the responsibility and resources for conducting analytic epidemiologic investigations of residents of communities in the vicinity of DOE facilities, workers at DOE facilities, and other persons potentially exposed to radiation or to potential hazards from non-nuclear energy production and use. HHS has delegated program responsibility to CDC. Community involvement is a critical part of ATSDR's and CDC's energy-related research and activities, and input from members of the ORRHES is part of these efforts.

Purpose: The purpose of this meeting is to address issues that are unique to community involvement with the ORRHES, and agency updates.

Matters To Be Discussed: Agenda item will include a discussion on the draft TSCA Public Health Assessment, comments from the Exposure Investigation Workgroup, and updates from the Agency.

Agenda items are subject to change as priorities dictate.

For Further Information Contact: Marilyn (Palmer) Horton, Designated Federal Official and Health Communications Specialist, Division of Health Assessment and Consultation, ATSDR, 1600 Clifton Road, NE M/S E-32 Atlanta, Georgia 30333, telephone 1-

888-42-ATSDR (28737), fax 404/498-1744.

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 ATSDR.

Dated: April 15, 2005.

Alvin Hall,

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

[FR Doc. 05-7994 Filed 4-20-05; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Administration on Children, Youth and Families 2005 Head Start Tribally Controlled Land Grant College and University Partnerships; Notice of Correction for the FY 05 Head Start Tribally Controlled Land Grant College and University Partnerships Program Announcement, HHS-2005-ACF-ACYF-YT-0012, CFDA# 93.600

AGENCY: Administration on Children, Youth and Families, Head Start Bureau, ACF, DHHS.

ACTION: Notice of corrections.

SUMMARY: This notice is to inform interested parties of corrections to the Head Start Tribally Controlled Land Grant College and University Partnerships Program Announcement that was published on Wednesday, April 13, 2005. The following corrections should be noted:

(1) Under Priority Areas I, Section VII. Agency Contacts, Program Office Contact, please delete the following name, address, phone number, and e-mail address: Katherine Gray, U.S. Department of Health and Human Services, Administration for Children and Families, ACYF—Head Start Bureau, 330 C Street, SW., Switzer Room 2211, Washington, DC 20447. Phone: 312-353-2260. E-mail: kgray@acf.hhs.gov.

Please replace the deleted name, address, phone number, and e-mail address with the following: Rosalind Dailey, U.S. Department of Health and Human Services, Administration for Children and Families, ACYF—Head Start Bureau, 330 C Street, SW., Switzer Room 2211, Washington, DC 20447. Phone: 202-205-8653. E-mail: rdailey@acf.hhs.gov.

All information in this notice of correction is accurate and replaces information specified in the April 13 notice. Applications are still due by the deadline date that was published in the April 13 notice (May 13 for Letters of Intent or Preapplications and June 13 for Applications).

FOR FURTHER INFORMATION CONTACT: For further information please contact the Administration on Children, Youth and Families, Head Start Bureau at (202) 205-8653 or rdailey@acf.hhs.gov.

Dated: April 14, 2005.

Joan E. Ohl,

Commissioner, Administration on Children, Youth and Families.

[FR Doc. 05-7949 Filed 4-20-05; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Anesthesiology and Respiratory Therapy 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: Anesthesiology and Respiratory Therapy 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 May 13, 2005, from 8 a.m. to 5:30 p.m.

Location: Hilton Washington DC North/Gaithersburg, Salons A and B, 620 Perry Pkwy., Gaithersburg, MD.

Contact Person: Neel Patel, Center for Devices and Radiological Health (HFZ-480), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301-443-8611, ext. 3, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 3014512624. Please call the Information Line for up-to-date information on this meeting.

Agenda: The committee will hear a presentation on FDA's Critical Path Initiative and a presentation by the Office of Surveillance and Biometrics in