(Pub. L. 92–463), the Centers for Disease Control and Prevention, NCEH/ATSDR announces the following subcommittee meeting:

Name: Community and Tribal Subcommittee (CTS).

Time and Date: 3 p.m.–4:30 p.m., September 8, 2005.

Place: The teleconference will originate at the National Center for Environmental Health/Agency for Toxic Substances and Disease Registry in Atlanta, Georgia. Please see *Supplementary Information* for details on accessing the teleconference.

Status: Open to the public, teleconference access limited only by availability of telephone ports.

Purpose: Under the charge of the Board of Scientific Counselors, NCEH/ATSDR the Community and Tribal Subcommittee will provide the BSC, NCEH/ATSDR with a forum for community and tribal first-hand perspectives on the interactions and impacts of the NCEH/ATSDR's national and regional policies, practices and programs.

Matters To Be Discussed: The teleconference agenda will include an update on the Report on the Program Peer Review Subcommittee, a discussion on the NCEH/ATSDR portfolio of programs; and an open discussion for other important issues.

Items are subject to change as priorities dictate.

Supplementary Information: This conference call is scheduled to begin at 3 p.m. eastern standard time. To participate in the teleconference, please dial (877) 315–6535 and enter conference code 383520.

For Further Information Contact: Sandra Malcom, Committee Management Specialist, Office of Science, NCEH/ATSDR, M/S E–28, 1600 Clifton Road, NE., Atlanta, Georgia 30333, telephone (404) 498–0003.

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 National Center for Environmental Health/Agency for Toxic Substances and Disease Registry.

Dated: August 25, 2005.

B. Kathy Skipper,

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

[FR Doc. 05–17295 Filed 8–30–05; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Agency for Toxic Substances and Disease Registry (ATSDR) Public Meeting of the Citizens Advisory Committee on Public Health Service (PHS) Activities and Research at Department of Energy (DOE) Sites: Oak Ridge Reservation Health Effects Subcommittee

Name: Public meeting of the Citizens Advisory Committee on PHS Activities and Research at DOE Sites: Oak Ridge Reservation Health Effects Subcommittee (ORRHES).

Time and Date: 12 p.m.—8 p.m., September 22, 2005.

Place: Oak Ridge Mall, Alpine Room, 333 East Main Street, Oak Ridge, Tennessee 37830.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 50 people.

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 ''Superfund''). 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 (DHHS) 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. DHHS 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 items will include a brief discussion on the Beir VII report; a presentation on the draft public health assessment: Current and Future Chemical Exposure Evaluation (1990–2003); an update on ATSDR's project management plan and the schedule of public health assessments to be released in FY2005–2006; updates and recommendations from the Exposure Evaluation, Community Concerns and Communications, and the Health Outcome Data Workgroups; and agency updates.

Agenda items are subject to change as priorities dictate.

For Further Information Contact: Marilyn Horton, Designated Federal Official and Health Communication 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 ATDSR.

Dated: August 25, 2005.

B. Kathy Skipper,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Educational Workshops on Current Good Manufacturing Practices; Public Workshops

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshops.

SUMMARY: The Food and Drug Administration (FDA) is announcing a series of educational workshops on current good manufacturing practice (CGMP). The workshops, which will be held in collaboration with Peking University (Beijing, China) and the International Society for Pharmaceutical Engineering (ISPE), are intended to educate participants on current methods for compliance with good manufacturing practices (GMP). The workshops are being offered to help ensure effective CGMP programs and to further the common goals of FDA and providers of quality pharmaceutical products.

DATES: See table 1 in the **SUPPLEMENTARY INFORMATION** section of this document.

ADDRESSES: See table 1 in the **SUPPLEMENTARY INFORMATION** section of this document.

FOR FURTHER INFORMATION CONTACT: Erik N. Henrikson, Center for Drug Evaluation and Research (HFD–320), Food and Drug Administration, 11919 Rockville Pike, Rockville, MD 20852,