Administration, 10903 New Hampshire Ave., Bldg. 31, rm. 2417, Silver Spring, MD 20993–0002, 301–769–9001, FAX: 301–847–8533, email: diem.ngo@fda.hhs.gov, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), codes 3014512543 and 3014512535. 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 November 3, 2010, the committees will discuss a number of safety concerns with intravenous administration of the anti-seizure drugs phenytoin and fosphenytoin, including the condition known as Purple Glove Syndrome, and recommend what regulatory actions, if any, are necessary to diminish the risks.

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 October 20, 2010. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. Those desiring to make 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 October 12, 2010. Time allotted for each presentation may be limited. If the number of registrants requesting greater time 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 October 13, 2010.

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 Diem-Kieu Ngo 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).


Leslie Kux,
Acting Assistant Commissioner for Policy.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Institute for Occupational Safety and Health (NIOSH), Safety and Occupational Health Study Section (SOHSS)

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 following committee meeting.

Times and Dates:
8 a.m.–5 p.m., October 7, 2010 (Closed);
8 a.m.–5 p.m., October 8, 2010 (Closed).
Place: Embassy Suites Hotel, 1900 Diagonal Road, Alexandria, Virginia, 22314, Telephone (703) 684–5900, Fax (703) 684–1403.
Purpose: The Safety and Occupational Health Study Section will review, discuss, and evaluate grant application(s) received in response to the Institute’s standard grants review and funding cycles pertaining to research issues in occupational safety and health, and allied areas.

It is the intent of NIOSH to support broad-based research endeavors in keeping with the Institute’s program goals. This will lead to improved understanding and appreciation for the magnitude of the aggregate health burden associated with occupational injuries and illnesses, as well as to support more focused research projects, which will lead to improvements in the delivery of occupational safety and health services, and the prevention of work-related injury and illness. It is anticipated that research funded will promote these program goals.

Matters to be Discussed: The meeting agenda includes discussions on matters related to the conduct of Study Section business and for the study section to consider safety and occupational health-related grant applications. These portions of 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, Centers for Disease Control and Prevention, pursuant to Section 10(d) Public Law 92–463.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information:
Price Connor, PhD, NIOSH Health Scientist, 1600 Clifton Road, NE., Mailstop E–20, Atlanta, Georgia 30333, Telephone (404) 498–2511, Fax (404) 498–2571.

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.


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

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Center for Health Statistics (NCHS), Classifications and Public Health Data Standards Staff, Announces the Following Meeting

Name: ICD–9–CM Coordination and Maintenance Committee meeting.
Time and Date: 9 a.m.–5:30 p.m., September 15–16, 2010.
Place: Centers for Medicare and Medicaid Services (CMS) Auditorium, 7500 Security Boulevard, Baltimore, Maryland.
Status: Open to the public.
Purpose: The ICD–9–CM Coordination and Maintenance (C&M) Committee will hold the last meeting of the 2010 calendar year cycle on Wednesday and Thursday September 15–16, 2010. The C&M meeting, is a public forum for the presentation of proposed modifications to the International Classification of Diseases, Ninth-Revision, Clinical Modification. There will be telephone lines available from 9 a.m. until 12:30 p.m. and 2:15 p.m. until 5:30 p.m. (Wednesday) and 9 until 12:30 and 1:30 until 3:15 p.m. (Thursday) for those who are unable to attend the meeting in person.
Tentative agenda items include:

September 15, 2010
- ICD–10 Topics (9–12:30)
- Freeze update
- General Equivalence Maps (GEMs)*
- MS–DRG Impact Analysis
- V28.0 ICD–10 MS–DRGs
- ICD–10–CM/ICD–10–PCS updates
- * Section 10109(c) of the Patient Protection and Affordable Care Act and the Reconciliation Act of 2010 (PPACA) requires the Secretary of Health and Human Services (HHS) to task the C&M Committee to convene a meeting before January 1, 2011, to receive stakeholder input regarding the crosswalk between the Ninth and Tenth Revisions of the International Classification of Diseases (ICD–9, and ICD–10, respectively), posted to the CMS Web site at http://www.cms.gov/ICD10, for the purpose of making appropriate revisions to said crosswalk.
- Section 10109(c) further states that any revised crosswalk be treated as a code set for which a standard has been adopted by the Secretary, and that revisions to this crosswalk be posted to the CMS Web site.
- The C&M Committee will use the first half of the first day of the September C&M Committee meeting, 9 a.m. to 12:30 p.m. Wednesday, September 15, 2010, to fulfill the above-referenced PPACA requirements for this meeting to be held prior to January 1, 2011, and receive public input regarding the above-referenced crosswalk revision.
- No other meeting will be convened by the C&M Committee for this purpose. Interested parties and stakeholders should be prepared to submit their written comments and other relevant documentation at the meeting, or no later than November 12, 2010 to the following addresses:
  - ICD–9–CM procedures:
    - Contrast Dye Removal
    - Endovascular partial occlusion of abdominal aorta
    - Endovascular Intracranial Aneurysm Embolization
    - Penetrated AAA Endovascular Graft Implantation of an Anti-Microbial Envelope
    - Pulmonary Artery Pressure Monitoring Addenda (procedures)
  - September 16, 2010
    - Corticobasal degeneration
    - Complication of stem cell transplant
    - Gastraparesis
    - Glaucoma
    - Hepatopulmonary syndrome
    - Intestinal lung diseases
    - Malnutrition
    - Mesh erosion
    - Pseudobulbar affect
    - Transfusion transmitted infections
    - Genitourinary conditions
    - Addenda (diagnoses)
  - Contact Person for Additional Information: Donna Pickett, Medical Systems Administrator, Classifications and Public Health Data Standards Staff, NCHS, 3311 Toledo Road, Room 2337, Hyattsville, Maryland 20782, e-mail dfp4@cdc.gov, telephone 301–458–4434 (diagnosis), Mady Hue, Health Insurance Specialist, Division of Acute Care, CMS, 7500 Security Blvd., Baltimore, Maryland 21244, e-mail marilu.hue@cms.hhs.gov, telephone 410–786–4510 (procedures).
  - Note: CMS and NCHS will no longer be providing paper copies of handouts for the meeting. Electronic copies of all meeting materials will be posted on the CMS and NCHS Web sites prior to the meeting at http://www.cms.hhs.gov/ICDPProviderDiagnosticCodes/03
    - Addenda (procedures)
    - Section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of meetings of the National Advisory Neurological Disorders and Stroke Council.
  - Name of Committee:
  - National Institute of Neurological Disorders and Stroke; Notice of Closed Meeting
  - Dated: September 8, 2010.
  - Elaine L. Baker,
  - Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.
  - [FR Doc. 2010–23092 Filed 9–15–10; 8:45 am]

OMB Control Number: 1056–0001
DEPARTMENT OF HEALTH AND HUMAN SERVICES
National Institutes of Health

National Institute of Neurological Disorders and Stroke; Notice of Closed Meeting
Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of meetings of the National Advisory Neurological Disorders and Stroke Council.
The meeting will be closed to the public as indicated below in accordance with the provisions set forth in sections 552b(c)(6). Title 5 U.S.C., as amended for the review, discussion, and evaluation of individual intramural programs and projects conducted by the National Institute of Neurological Disorders and Stroke, including consideration of personnel qualifications and performance, and the competence of individual investigators, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.
Name of Committee: Board of Scientific Counselors, National Institute of Neurological Disorders and Stroke.
Date: October 24–25, 2010.
Time: 7 p.m. to 6:30 p.m.
Agenda: To review and evaluate personal qualifications and performance, and competence of individual investigators.
Place: Hyatt Regency Bethesda, One Bethesda Metro Center, 7406 Wisconsin