

Dated: July 24, 2008.

**Penelope Slade Royall,**

*RADM, USPHS, Deputy Assistant Secretary for Health, (Disease Prevention and Health Promotion).*

[FR Doc. E8-18299 Filed 8-7-08; 8:45 am]

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

**National Center for Preparedness, Detection, and Control of Infectious Diseases**

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.

*Name:* Clinical Laboratory Improvement Advisory Committee (CLIAC).

*Times and Dates:* 8:30 a.m.–5 p.m., September 10, 2008; 8:30 a.m.–3 p.m., September 11, 2008.

*Place:* Centers for Disease Control and Prevention, 1600 Clifton Road, NE., Tom Harkin Global Communications Center, Building 19, Room 232, Auditorium B, Atlanta, Georgia 30333.

*New Information—Online Registration Required:* In order to expedite security clearance process at the CDC Roybal Campus located on Clifton Road, all CLIAC attendees are required to register in advance for the meeting at <http://www.cdc.gov/cliac/default.aspx> by clicking the Register for a "Meeting" link and completing all forms according to the instructions given. Please complete all the required fields and submit your registration as far in advance of the meeting date as possible.

**Note:** The cut-off date for registration for domestic attendees is Thursday, September 4, 2008; the cut-off date for international attendees to register is Monday, August 25, 2008.

*Status:* Open to the public, limited only by the space available. The meeting Room accommodates approximately 100 people.

*Purpose:* This Committee is charged with providing scientific and technical advice and guidance to the Secretary of Health and Human Services, the Assistant Secretary for Health, and the Director, CDC, regarding the need for, and the nature of, revisions to the standards under which clinical laboratories are regulated; the impact on medical and laboratory practice of proposed revisions to the standards; and the modification of the standards to accommodate technological advances.

*Matters to Be Discussed:* The agenda will include updates from the CDC, the Centers for Medicare & Medicaid Services, and the Food and Drug Administration; a report from the CLIAC Workgroup on Good Laboratory Practices for Genetic Testing, and discussion

of the Workgroup's proposals related to such: presentations and discussion related to laboratory quality control through risk management; and an introduction to the status of waived testing and discussion of the potential for waiver of automated hematology devices. Agenda items are subject to change as priorities dictate.

*Providing Oral or Written Comments:* It is the policy of CLIAC to accept written public comments and provide a brief period for oral public comments whenever possible.

*Oral Comments:* In general, each individual or group requesting to make an oral presentation will be limited to a total time of five minutes (unless otherwise indicated). Speakers must also submit their comments in writing for inclusion in the meeting's Summary Report. To assure adequate time is scheduled for public comments, individuals or groups planning to make an oral presentation should, when possible, notify the contact person below at least one week prior to the meeting date.

*Written Comments:* For individuals or groups unable to attend the meeting, CLIAC accepts written comments until the date of the meeting (unless otherwise stated). However, the comments should be received at least one week prior to the meeting date so that the comments may be made available to the Committee for their consideration and public distribution. Written comments, one hard copy with original signature, should be provided to the contact person below. Written comments will be included in the meeting's Summary Report.

*Contact Person for Additional Information:* Nancy Anderson, Chief, Laboratory Practice Standards Branch, Division of Laboratory Systems, National Center for Preparedness, Detection, and Control of Infectious Diseases, Coordinating Center for Infectious Diseases, CDC, 1600 Clifton Road, NE., Mailstop F-11, Atlanta, Georgia 30333; telephone (404) 498-2741; fax (404) 498-2219; or via e-mail at [Nancy.Anderson@cdc.hhs.gov](mailto:Nancy.Anderson@cdc.hhs.gov).

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 CDC and the Agency for Toxic Substances and Disease Registry.

Dated: July 28, 2008.

**Elaine L. Baker,**

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

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**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; Modifications to the International Classification of Diseases, Ninth Revision, Clinical Modification (ICD-9-CM), Supplementary Classification of External Causes of Injury and Poisoning**

**ACTION:** Notice.

National Center for Health Statistics (NCHS), Classifications and Public Health Data Standards Staff, announces the following modifications to the International Classification of Diseases, Ninth Revision, Clinical Modification (ICD-9-CM), Supplementary Classification of External Causes of Injury and Poisoning. These codes will become effective October 1, 2008.

**External Cause Tabular**

New code E927.0 Overexertion from sudden strenuous movement. Sudden trauma from strenuous movement

New code E927.1 Overexertion from prolonged static position

New code E927.2 Excessive physical exertion from prolonged activity

New code E927.3 Cumulative trauma from repetitive motion

New code E927.4 Cumulative trauma from repetitive impact

New code E927.8 Other overexertion and strenuous and repetitive movements or loads

New code E927.9 Unspecified overexertion and strenuous and repetitive movements or loads

**CONTACT PERSON FOR ADDITIONAL**

**INFORMATION:** Donna Pickett, Medical Systems Specialist, Classifications and Public Health Data Standards Staff, NCHS, 3311 Toledo Road, Room 2402, Hyattsville, Maryland 20782, e-mail [djp4@cdc.gov](mailto:djp4@cdc.gov), telephone 301-458-4434. The complete diagnosis addenda may be accessed on the NCHS Web site using the URL: <http://www.cdc.gov/nchs/datawh/ftperv/ftp9/ftp9.htm#addenda>.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to modifications to the ICD-9-CM, for both CDC and the