

Dated: February 25, 2016.

Lorin S. Curit,

*Director, Federal Acquisition Policy Division,
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Governmentwide Policy.*

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Initial Review

The meeting announced below concerns Childhood Obesity Research Demonstration 2.0, FOA DP 16-004, initial review.

SUMMARY: This document corrects a notice that was published in the **Federal Register** on February 10, 2016, Volume 81, Number 27, pages 7123-7124. The meeting time and date should read as follows:

Time and Date: 10:00 a.m.–6:00 p.m., EDT, March 15, 2016 (Closed).

FOR FURTHER INFORMATION CONTACT: Jaya Raman, Ph.D., Scientific Review Officer, CDC, 4770 Buford Highway NE., Mailstop F80, Atlanta, Georgia 30341, Telephone: (770) 488-6511, KVA5@cdc.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 both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

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

Clinical Laboratory Improvement Advisory Committee

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: 8:30 a.m.–5:00 p.m., April 13, 2016; 8:30 a.m.–12:00 p.m., April 14, 2016.

Place: CDC, 1600 Clifton Road NE., Tom Harkin Global Communications Center, Building 19, Auditorium B, Atlanta, Georgia 30333.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 100 people. This meeting will also be webcast, please see information below.

Purpose: This Committee is charged with providing scientific and technical advice and guidance to the Secretary of Health and Human Services (HHS); the Assistant Secretary for Health; the Director, Centers for Disease Control and Prevention; the Commissioner, Food and Drug Administration (FDA); and the Administrator, Centers for Medicare and Medicaid Services (CMS). The advice and guidance pertain to general issues related to improvement in clinical laboratory quality and laboratory medicine practice and specific questions related to possible revision of the Clinical Laboratory Improvement Amendment (CLIA) standards. Examples include providing guidance on studies designed to improve safety, effectiveness, efficiency, timeliness, equity, and patient-centeredness of laboratory services; revisions to the standards under which clinical laboratories are regulated; the impact of proposed revisions to the standards on medical and laboratory practice; and the modification of the standards and provision of non-regulatory guidelines to accommodate technological advances, such as new test methods, the electronic transmission of laboratory information, and mechanisms to improve the integration of public health and clinical laboratory practices.

Matters For Discussion: The agenda will include agency updates from CDC, CMS, and FDA. Presentations and discussions will include methods for improving the effectiveness/efficiency of CLIA meetings; an overview of the CMS Advisory Panel on Clinical Diagnostic Laboratory Tests; laboratory interoperability including the Office of the National Coordinator for Health Information Technology (ONC) policies and engagement with clinical laboratories; update on the cytology workload project; update on laboratory biosafety in clinical laboratories; and future CLIA topics.

Agenda items are subject to change as priorities dictate.

Webcast: The meeting will also be webcast. Persons interested in viewing

the webcast can access information at: <http://cdclabtraining.adobeconnect.com/aprilcliac/>.

In-Person Attendance Online Registration Required: All people attending the CLIA meeting in-person are required to register for the meeting online at least 5 business days in advance for U.S. citizens and at least 10 business days in advance for international registrants. Register at: <http://www.cdc.gov/cliac/Meetings/MeetingDetails.aspx#>.

Register by scrolling down and clicking the "Register for this Meeting" button and completing all forms according to the instructions given. Please complete all the required fields before submitting your registration and submit no later than April 7, 2016 for U.S. registrants and March 31, 2016 for international registrants.

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

Oral Comments: In general, each individual or group requesting to make oral comments 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, speakers should 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, CLIA accepts written comments until the date of the meeting (unless otherwise stated). However, it is requested that comments be submitted at least one week prior to the meeting date so that the comments may be made available to the Committee for their consideration and for public distribution. Written comments, one hard copy with original signature, should be provided to the contact person listed below, and will be included in the meeting's Summary Report.

Availability of Meeting Materials: To support the green initiatives of the federal government, the CLIA meeting materials will be made available to the Committee and the public in electronic format (PDF) on the internet instead of by printed copy. Check the CLIA Web site on the day of the meeting for materials: http://wwwn.cdc.gov/cliac/cliac_meeting_all_documents.aspx.

Note: If using a mobile device to access the materials, please verify that the device's browser is able to download the files from the CDC's Web site before the meeting. Alternatively, the files can be downloaded to