

choose voluntarily to be subject to section 4205.

- Factors to consider with respect to availability and use of space on menus and menu boards.

B. Determination of Calorie Content of Foods Offered by Chain Retail Food Establishments

- Information about standardization of recipes and methods of preparation.
- Information about variation in serving size and formulation of menu items.
- The role of inadvertent human error.
- Information about training of food service workers.
- Information about variations in ingredients.
- Any other relevant factors.

C. Vending Machine Operations

- Current practices within the vending machine industry with respect to the availability to prospective purchasers of Nutrition Facts panel information or otherwise providing visible nutrition information at the point of purchase.
- Possible mechanisms for displaying products' Nutrition Facts panels or otherwise providing visible nutrition information at the point of purchase.
- Factors to consider with respect to availability and use of space on vending machines.
- Considerations in requiring caloric content disclosure about food items sold from vending machines.
- Information about the size of chain vending machine operators (e.g., based on annual revenue or on number of locations).
- Information about the number of chain vending machine operators that are or could be affected by section 4205.
- Information about vending machine operators not covered by section 4205 of the Affordable Care Act that might elect to be subject to the requirements of section 4205.

D. Implementation and Enforcement

- Information about implementation, including information about options for inspection and enforcement.
- Information about inspection and enforcement mechanisms in state and local nutrition labeling programs.

III. Comments

Interested persons may submit to the Division of Dockets Management (see **ADDRESSES**) either electronic or written comments regarding this document. It is only necessary to send one set of comments. It is no longer necessary to send two copies of mailed comments.

Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

IV. References

1. Howlett, E.A., S. Burton, K. Bates, and K. Huggins, "Coming to a Restaurant Near You? Potential Consumer Responses to Nutrition Information Disclosure on Menus," Available at <http://www.journals.uchicago.edu/doi/pdf/10.1086/598799> *Journal of Consumer Research* 36(3): 494–503, 2009.
2. Tandon, P.S., J. Wright, C. Zhou, C.B. Rogers, D.A. Christakis, "Nutrition Menu Labeling May Lead to Lower-Calorie Restaurant Meal Choices for Children," *Pediatrics* 125(2): 244–248, 2010.
3. Center for Science in the Public Interest, "Comparison of Menu Labeling Policies," Available at http://cspinet.org/new/pdf/comparison_of_ml_policies_6-9.pdf.

Dated: June 29, 2010.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2010–16303 Filed 7–6–10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Clinical Laboratory Improvement Advisory Committee (CLIAC)

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 meeting of the aforementioned committee:

Times and Dates:

8:30 a.m.–5 p.m., September 1, 2010.

8:30 a.m.–3 p.m., September 2, 2010.

Place: CDC, 1600 Clifton Road, NE., Tom Harkin Global Communications Center, Building 19, Room 232, Auditorium B, Atlanta, Georgia 30333.

Online Registration Required: In order to expedite the security clearance process at the CDC Roybal Campus located on Clifton Road, all CLIAC attendees are required to register for the meeting online at least 14 days in advance at <http://wwwn.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 before submitting your registration and submit no later than August 17, 2010.

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 agency updates from the CDC, the Centers for Medicare & Medicaid Services, and the Food and Drug Administration. Reports and Committee discussion will address issues pertaining to cytology testing and workload recording; the electronic exchange of laboratory information; and consideration of proposals from the CLIAC proficiency testing workgroup. 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 Science and Standards (proposed), Laboratory Science, Policy and Practice Program Office (LSPPO) (proposed), Office of Surveillance, Epidemiology and Laboratory Services (proposed), Centers for Disease Control and Prevention, 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.

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: June 28, 2010.

Elaine L. Baker,

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

[FR Doc. 2010–16387 Filed 7–6–10; 8:45 am]

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