DEPARTMENT OF HEALTH AND HUMAN SERVICES
Centers for Disease Control and Prevention
Advisory Committee to the Director (ACD), Centers for Disease Control and Prevention—National Biosurveillance Advisory Subcommittee (NBAS)

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 aforementioned subcommittee:

Time and Date: 8:30 a.m.–11:30 a.m., March 21, 2011.

Place: Emory Conference Center Hotel, 1615 Clifton Road, NE., Atlanta, Georgia 30329. Telephone: (404) 712–6000.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 75 people. The public is welcome to participate during the public comment periods. The public comment period is tentatively scheduled for 11 a.m.–11:15 a.m.

Purpose: As a subcommittee to the CDC’s Advisory Committee to the Director (ACD), the NBAS will provide counsel to the ACD and the Federal government through the ACD regarding a broad range of human health surveillance issues arising from the development and implementation of a roadmap for the human health component of a national biosurveillance system.

Matters to be Discussed: Agenda items will include the subcommittee’s discussion, deliberation, and vote on the proposed report for enhancing the nation’s biosurveillance capability. The agenda is subject to change as priorities dictate.

Contact Person for More Information:
Pamela Diaz, M.D., Designated Federal Officer, ACD, CDC—NBAS, 1600 Clifton Road, NE., M/S E–97, Atlanta, Georgia 30333. Telephone: (404) 498–0476. E-mail: pdiaz@cdc.gov. For security reasons, members of the public interested in attending the meeting should contact Mark Byers. Telephone: (404) 498–0481. E-mail: mbyers@cdc.gov. The deadline for notification of attendance is March 10, 2011.

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.

Dated: February 25, 2011.

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

Correction: This notice was published in the Federal Register on January 21, 2011, Volume 76, Number 14, Page 3908. The date for the aforementioned meeting has been changed to the following:

DATES: April 26, 2011 (Closed)

Contact Person for More Information:
Michael Dalmat, Dr.P.H., Scientific Review Officer, CDC, National Center for Chronic Disease Prevention and Health Promotion, Office of the Director, Extramural Research Program Office, 4770 Buford Highway, NE., Mailstop K–92, Atlanta, Georgia 30341. Telephone: (770) 488–6423. E-mail: MED1@CDC.GOV.

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Dated: February 25, 2011.

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

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Centers for Medicare & Medicaid Services

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare & Medicaid Services

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare & Medicaid Services (CMS), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the Agency’s function; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

1. Type of Information Collection Request: Extension without change of a currently approved collection; Title of Information Collection: Clinical Laboratory Improvement Amendments of 1988 (CLIA) Budget Workload Reports and Supporting Regulations in 42 CFR 493.1–2001; Use: The collected information will be used by CMS to determine the amount of Federal reimbursement for surveys conducted. Use of the information includes program evaluation, audit, budget formulation and budget approval. Form CMS–102 is a multi-purpose form designed to capture and record all budget and expenditure data. Form CMS–105 captures the annual projected CLIA workload that the State survey agency will accomplish. It is also used by the CMS regional office to approve the annual projected CLIA workload. The information is required as part of the section 1864 agreement with the State; Form Numbers: CMS–102 and CMS–105 (OMB#: 0938–0509); Frequency: Quarterly. Affected Public: State, Local, or Tribal Governments: Number of Respondents: 50; Total Annual Responses: 50; Total Annual Hours: 4,500. (For policy questions regarding