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 Human Immunodeficiency Virus (HIV) Prevention Projects for Young Men of Color Who Have Sex with Men and Young Transgender Persons of Color, Funding Opportunity Announcement (FOA) PS11–1113, initial review.

Correction: The notice was published in the Federal Register on July 7, 2011, Volume 76, Number 130, Page 39879. The place should read as follows:

Place: Hilton Atlanta Hotel, 255 Courtland Street, NE., Atlanta, Georgia 30303. Telephone: (404) 498–2726.

FOR FURTHER INFORMATION CONTACT:

Harriette Lynch, Public Health Analyst, Extramural Programs, National Center for HIV, Hepatitis and Sexually Transmitted Diseases Prevention, CDC, 1600 Clifton Road, NE., Mailstop E–60, Atlanta, Georgia 30333. Telephone: (404)498–2726, E-mail: HLynch@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.

Dated: July 22, 2011.

Elaine L. Baker,

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

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 of a currently approved collection; Title of Information Collection: Consumer Assistance Program Grants; Use: Section 1002 of the Affordable Care Act provides for the establishment of consumer assistance (or ombudsman) programs, starting in FY 2010. Federal grants will support these programs. For FY 2010, $30 million is appropriated. These programs will assist consumers with filing complaints and appeals, assist consumers with enrollment into health coverage, collect data on consumer inquiries and complaints to identify problems in the marketplace, educate consumers on their rights and responsibilities, and starting in 2014, resolve problems with premium credits for Exchange coverage. Importantly, these programs must provide detailed reporting on the types of problems and questions consumers may experience with health coverage, and how these are resolved. In order to strengthen oversight, the law requires programs to report data to the Secretary of the Department of Health and Human Services (HHS) “As a condition of receiving a grant under subsection (a), an office of health insurance consumer assistance or ombudsman program shall be required to collect and report data to the Secretary on the types of problems and inquiries encountered by consumers” (Sec. 2793 (d)). Form Number: CMS–10333 (OMB–0938–1097); Frequency: Quarterly; Affected Public: Private Sector; State, Local, or Tribal Governments; Number of Respondents: 40; Number of Responses: 200; Total Annual Hours: 4,800. [For policy questions regarding this collection, contact Eliza Bangit at (301) 492–4219. For all other issues call (410) 786–1326.]

2. Type of Information Collection Request: New Collection; Title of Information Collection: Health Insurance Assistance Database; Use: In October 2010, the Office of Consumer Support began to take and respond to direct consumer inquiries related to the Affordable Care Act. As of February 15th 2011, CCIIO has received 906 consumer inquiries. Consumer inquiries continue to come in to CCIIO at a rate of 30 to 35 inquiries per week. Starting in January 2011, the HHS Hotline will begin to refer ACA calls to CCIIO. To date, the HHS Hotline receives, on average, 400 calls per month pertaining to ACA.

Accordingly, a system to collect, track and store consumer information is urgently needed in order to accomplish successful case management to ensure that the information, coverage, and health care needs of consumers are addressed fairly and in a timely fashion. Further, the Team will provide detailed reports on these consumer inquiries with a focus on Affordable Care Act and PHS Act compliance issues. These reports will assist the Office of Oversight in identifying areas where compliance concerns may arise. Reports will be stripped of any information in identifiable form (IIF) and personal health information when written and prepared. Authority for maintenance, collection and disclosures of this information is given under sections 2719, 2723, and 2761 of the Public Health Service Act (PHS Act) and section 1321(c) of the Affordable Care Act.

Analysis of this data reporting will help identify patterns of practice in the insurance marketplaces and uncover suspected patterns of noncompliance. HHS may share program data reports with the Departments of Labor and Treasury, and State regulators. Program data also can offer CCIIO one indication of the effectiveness of State enforcement, affording opportunities to provide technical assistance and support to State insurance regulators and, in extreme cases, inform the need to trigger Federal enforcement. Form Number: CMS–10384 (OGN: 0938–New); Frequency: Occasionally;