attending, and if participating via webcast or in-person.

Webcast Public Participation: After pre-registration, individuals participating by webcast will receive webcast access information via email.

In-Person Public Participation and Building Access: For in-person participants, the meetings are within the National Institutes of Health (NIH) Clinical Center (Building 10) as noted above in the Addresses section. Details regarding registration capacity and directions will be posted on *www.DietaryGuidelines.gov.* For in-person participants, check-in at the registration desk onsite at the meeting is required and will begin at 7:30 a.m. each day.

Public Comments and Meeting Documents: Written comments from the public will be accepted throughout the Committee's deliberative process; opportunities to present oral comments to the Committee will be provided at a future meeting. Written public comments can be submitted and/or viewed at *www.DietaryGuidelines.gov* using the ''Submit Comments'' and ''Read Comments'' links, respectively. Written comments received by June 5, 2013 will ensure transmission to the Committee prior to this meeting. Documents pertaining to Committee deliberations, including meeting agendas, summaries, and transcripts will be available on *www.DietaryGuidelines.gov* under ''Meetings'' and meeting materials will be available for public viewing at the meeting. Meeting information, thereafter, will continue to be accessible online, at the NIH Library, and upon request at the Office of Disease Prevention and Health Promotion, OASH/HHS; 1101 Wootton Parkway, Suite LL100 Tower Building; Rockville, MD 20852: Telephone (240) 453–8280; Fax: (240) 453–8281.

Dated: May 24, 2013.

Richard Olson,

Designated Federal Officer, Director, Division of Prevention Science, Office of Disease Prevention and Health Promotion, Office of the Assistant Secretary for Health, U.S. Department of Health and Human Services.

[FR Doc. 2013–12859 Filed 5–24–13; 4:15 pm]

BILLING CODE 4150–32–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Health Resources and Services Administration

CDC/HRSA Advisory Committee on HIV, Viral Hepatitis and STD Prevention and Treatment

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) and the Health Resources and Services Administration (HRSA) announce the following meeting of the aforementioned committee:

Times and Dates: 8:00 a.m.–5:30 p.m., June 18, 2013; 8:00 a.m.–2:30 p.m., June 19, 2013.

Place: CDC Corporate Square, Building 8, Conference Room 1–ABC, 8 Corporate Boulevard, Atlanta, Georgia 30329, Telephone: (404) 639–8317.

Status: Open to the public, limited only by the space available. The meeting room will accommodate approximately 100 people. This meeting is also accessible by teleconference. Toll-free +1 (866) 718–4584, Participant code: 8484551.

Status: Open to the public, limited only by the space available. The meeting room will accommodate approximately 100 people.

Purpose: This Committee is charged with advising the Director, CDC and the Administrator, HRSA, regarding activities related to prevention and control of HIV/AIDS, Viral Hepatitis and other STDs, the support of health care services to persons living with HIV/AIDS, and education of health professionals and the public about HIV/AIDS, Viral Hepatitis and other STDs.

Matters To Be Discussed: Agenda items include: (1) STD clinical preventive services in primary care setting and integrating STD screening and treatment services in HIV care settings); (2) The test and cure era for hepatitis C: The public health response to rising hepatitis C mortality; The impact of new therapies on health outcomes; and Building care capacity to increase access to hepatitis C virus (HCV) therapy; (3) HIV Medical Monitoring Project: follow up on Institute of Medicine (IOM) report and other Affordable Care Act (ACA) issues; (4) Recommendations for new HIV diagnostic laboratory testing algorithms; and (5) CHAC Workgroups Update.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Margie Scott-Cseh, Centers for Disease Control and Prevention, National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention, 1600 Clifton Road NE., Mailstop E–07, Atlanta, Georgia 30333, Telephone (404) 639–8317.

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.

[FR Doc. 2013–12857 Filed 5–29–13; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–0577]

Agency Information Collection Activities; Proposed Collection; Comment Request; Requirements for Submission of Labeling for Human Prescription Drugs and Biologics in Electronic Format

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection contained in the requirements for the submission of labeling for human prescription drugs and biologics in electronic format.

DATES: Submit either electronic or written comments on the collection of information by July 29, 2013.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.regulations.gov.* Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Ila S. Mizrachi, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., P150–400B, Rockville, MD 20850, 301–796–7726, *Ila.mizrachi@fda.hhs.gov.*