National Coordinator for review and coordination in the Eligibility/Enrollment Systems APD approval assignment. The information requested on the Checklist will be used to determine and approve enhanced FFP to States and to determine how States are complying with the seven standards and conditions; Form Number: CMS–10385 (OMB#: 0938–1125); Frequency: Occasionally; Number of Respondents: 56; Total Annual Responses: 168; Total Annual Hours: 204. (For policy questions regarding this collection contact Richard Friedman at 410–786–4451. For all other issues call 410–786–1326.)

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS Web Site address at http://www.cms.hhs.gov/PaperworkReductionActof1995, or E-mail your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@cms.hhs.gov, or call the Reports Clearance Office on (410) 786–1326.

To be assured consideration, comments and recommendations for the proposed information collections must be received by the OMB desk officer at the address below, no later than 5 p.m. on August 1, 2011.

OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer, Fax Number: (202) 395–6974, E-mail: OIRA_submission@omb.eop.gov.

Dated: June 28, 2011.

Michelle Shortt,
Director, Regulations Development Group, Office of Strategic Operations and Regulatory Affairs.

AGENCY: President’s Committee for People with Intellectual Disabilities (PCPID), HHS.

ACTION: Notice of committee meeting via conference call.

DATES: Tuesday, July 19, 2011, from 1 p.m. to 2:30 p.m. EST. This meeting, to be held via audio conference call, is open to the public.

Details for accessing the full Committee Conference Call are cited below: Toll Free Dial-In Number: 800–779–1436. Pass Code: PCPID.

Individuals who will need accommodations for a disability in order to participate in the PCPID Meeting via audio conferencing (assistive listening devices, materials in alternative format such as large print or Braille) should notify Genevieve Swift, PCPID Executive Administrative Assistant, at Edith.Swift@acf.hhs.gov, or by telephone at 202–619–0634, no later than Tuesday, July 12, 2011. PCPID will attempt to meet requests for accommodations made after that date, but cannot guarantee ability to grant requests received after this deadline.

AGENDA: Committee Members will discuss the potential topics, themes, and trends for the PCPID 2011 Annual Report to the President.


SUPPLEMENTARY INFORMATION: PCPID acts in an advisory capacity to the President and the Secretary of Health and Human Services, through the Administration on Developmental Disabilities, on a broad range of topics relating to programs, services, and supports for persons with intellectual disabilities. The PCPID Executive Order stipulates that the Committee shall: (1) Provide such advice concerning intellectual disabilities as the President or the Secretary of Health and Human Services may request; and (2) provide advice to the President concerning the following for people with intellectual disabilities: (A) Expansion of educational opportunities; (B) promotion of homeownership; (C) assurance of workplace integration; (D) improvement of transportation options; (E) expansion of full access to community living; and (F) increasing access to assistive and universally designed technologies.

Dated: June 27, 2011.

Laverdia Taylor Roach, PCPID.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Experimental Study of Format Variations in the Brief Summary of Direct-to-Consumer Print Advertisements

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by August 1, 2011.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–7285, or e-mailed to oira_submission@omb.eop.gov. All comments should be identified with the OMB control number 0910–New and title, “Experimental Study of Format Variations in the Brief Summary of Direct-to-Consumer Print Advertisements.” Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Elizabeth Berbakos, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., P150–400B, Rockville, MD 20850, 301–796–3792, Elizabeth.Berbakos@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Experimental Study of Format Variations in the Brief Summary of Direct-to-Consumer Print Advertisements—OMB Control Number 0910–New

Section 502(n) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 352(n)) specifies that ads for prescription drugs and biological products must provide a true statement of information “in brief summary”