PLACE: Federal Trade Commission Building, Room 532, 600 Pennsylvania Avenue, NW., Washington, DC 20580.

STATUS: Part of this meeting will be open to the public. The rest of the meeting will be closed to the public.

MATTERS TO BE CONSIDERED:
(1) Oral Argument in The North Carolina Board of Dental Examiners, Docket 9343
(2) Executive Session to follow Oral Argument in The North Carolina Board of Dental Examiners, Docket 9343.

CONTACT PERSON FOR MORE INFORMATION: Mitch Katz, FTC, Office of Public Affairs, 600 Pennsylvania Avenue, NW., Washington, DC 20580, (202) 326–2180; Recorded Message: (202) 326–2711

By direction of the Commission.

Donald S. Clark,
Secretary.

[FR Doc. 2011–27261 Filed 10–21–11; 8:45 am]

BILLING CODE 6750–01–M

GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–0163; Docket 2011–0079; Sequence 5]

Submission for OMB Review; General Services Administration; Information Specific to a Contract or Contracting Action; (Not Required by Regulation)

AGENCY: Office of the Chief Acquisition Officer, GSA.

ACTION: Notice of request for comments regarding a renewal to an existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a previously approved information collection requirement regarding information specific to a contract or contracting action (not required by regulation). A notice was published in the Federal Register at 76 FR 38396, on June 30, 2011. No comments were received.

Public comments are particularly invited on: Whether this collection of information is necessary and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate and based on valid assumptions and methodology; and ways to enhance the quality, utility, and clarity of the information to be collected.

DATES: Submit comments on or before: November 23, 2011.

FOR FURTHER INFORMATION CONTACT: William Clark, Procurement Analyst, Acquisition Policy Division, at telephone (202) 219–1813 or e-mail william.clark@gsa.gov.

ADDRESSES: Submit comments identified by Information Collection 3090–0163, Information Specific to a Contract or Contracting Action (Not Required by Regulation), by any of the following methods:
- Regulations.gov: http://www.regulations.gov. Submit comments via the Federal eRulemaking portal by inputting “Information Collection 3090–0163, Information Specific to a Contract or Contracting Action (Not Required by Regulation),” under the heading “Enter Keyword or ID” and selecting “Search.” Select the link “Submit a Comment” that corresponds with “Information Collection 3090–0163, Information Specific to a Contract or Contracting Action (Not Required by Regulation).” Follow the instructions provided at the “Submit a Comment” screen. Please include your name, company name (if any), and “Information Collection 3090–0163, Information Specific to a Contract or Contracting Action (Not Required by Regulation),” on your attached document.

Instructions: Please submit comments only and cite Information Collection 3090–0163, Information Specific to a Contract or Contracting Action (Not Required by Regulation), in all correspondence related to this collection. All comments received will be posted without change to http://www.regulations.gov, including any personal and/or business confidential information provided.

SUPPLEMENTARY INFORMATION:

A. Purpose
The General Services Administration (GSA) has various mission responsibilities related to the acquisition and provision of supplies, transportation, information technology, telecommunications, real property management, and disposal of real and personal property. These mission responsibilities generate requirements that are realized through the solicitation and award of public contracts. Individual solicitations and resulting contracts may impose unique information collection/reporting requirements on contractors, not required by regulation, but necessary to evaluate particular program accomplishments and measure success in meeting special program objectives.

B. Annual Reporting Burden
Respondents: 126,870.
Responses per Respondent: 1.35.
Total Responses: 171,275.
Hours per Response: 40.
Total Burden Hours: 68,510.
Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat (MVCB), 1275 First Street, NE., Washington, DC 20417, telephone (202) 501–4755. Please cite OMB Control No. 3090–0163, Information Specific to a Contract or Contracting Action (Not Required by Regulation), in all correspondence.

Dated: October 18, 2011.

Laura Auletta,
Acting Director, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy.

[FR Doc. 2011–27440 Filed 10–21–11; 8:45 am]

BILLING CODE 6820–61–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention (CDC)

Advisory Committee on Childhood Lead Poisoning Prevention (ACCLPP)

Correction: This notice was published in the Federal Register on October 6, 2011, Volume 76, Number 194, page 62071. Contact information changed to delete Nikki Walker’s name and telephone number and replaced with Tiffany Turner’s name and telephone number.

CONTACT PERSONS FOR MORE INFORMATION: Claudine Johnson, Program Operation Assistant or Tiffany Turner, Healthy Homes and Lead Poisoning Prevention Branch, Division of Environmental Emergency Health Services, NCEH, CDC, 4770 Buford Highway, NE., Mailstop F–60, Atlanta, Georgia 30341, telephone (770) 488–3629; Tiffany Turner (770) 488–0554; fax (770) 488–3635.

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
Announces the following meeting for the Board of Scientific Counselors, National Institute for Occupational Safety and Health (BSC, NIOSH). 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 for the aforementioned committee:

**Time and Dates:** 8:30 a.m.–3:15 p.m., November 15, 2011.

**Place:** Patriots Plaza I, 395 E. Street, SW., Room 9000, Washington, DC 20201.

**Status:** Open to the public, limited only by the space available. The meeting room accommodates approximately 50 people. This meeting is available by teleconference. Please dial (877) 328–2816 and enter code 6538291.

**Purpose:** The Secretary, the Assistant Secretary for Health, and by delegation the Director, Centers for Disease Control and Prevention, are authorized under Sections 301 and 308 of the Public Health Service Act to conduct directly or by grants or contracts, research, experiments, and demonstrations relating to occupational safety and health and to mine health. The Board of Scientific Counselors shall provide guidance to the Director, National Institute for Occupational Safety and Health on research and prevention programs. Specifically, the Board shall provide guidance on the Institute’s research activities related to developing and evaluating hypotheses, systematically documenting findings and disseminating results. The Board shall evaluate the degree to which the activities of the National Institute for Occupational Safety and Health: (1) Conform to appropriate scientific standards; (2) address current, relevant needs; and (3) produce intended results.

**Matters To Be Discussed:** The agenda will include the following: (1) Director Update; (2) Implementation of the National Academies Program Recommendations for Respiratory Diseases, Hearing Loss Prevention, Personal Protective Technology, and Health Hazard Evaluations; (3) Occupational Safety and Health Workforce Needs Assessment; (4) and Future Directions for Extramural Research. Agenda items are subject to change as priorities dictate.

**Contact Person for More Information:** Roger Rosa, Designated Federal Officer, BSC, NIOSH, CDC, 395 E Street, SW., Suite 9200, Patriots Plaza Building, Washington, DC 20201, telephone (202) 245–0655, fax (202) 245–0664.

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:** October 17, 2011.

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

**BILLING CODE 4163–18–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

**Board of Scientific Counselors, National Institute for Occupational Safety and Health (BSC, NIOSH)**

**Food and Drug Administration**

[Docket No. FDA–2011–N–0264]

**Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Request for Designation as Country Not Subject to the Restrictions Applicable to Human Food and Cosmetics Manufactured From, Processed With, or Otherwise Containing, Material From Cattle**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that a collection of information entitled “Request for Designation as Country Not Subject to the Restrictions Applicable to Human Food and Cosmetics Manufactured From, Processed With, or Otherwise Containing, Material From Cattle” has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

**FOR FURTHER INFORMATION CONTACT:** Denver Presley, Jr., Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., P150–400B, Rockville, MD 20850, 301–796–3793.

**SUPPLEMENTARY INFORMATION:** On June 28, 2011, the Agency submitted a proposed collection of information entitled “Request for Designation as Country Not Subject to the Restrictions Applicable to Human Food and Cosmetics Manufactured From, Processed With, or Otherwise Containing, Material From Cattle” to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0623. The approval expires on September 30, 2014. A copy of the supporting statement for this information collection is available on the Internet at http://www.reginfo.gov/public/do/PRAMain.

**Dated:** October 17, 2011.

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

**BILLING CODE 4163–18–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA–2011–N–0424]

**Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Temporary Marketing Permit Applications**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that a collection of information entitled “Temporary Marketing Permit Applications” has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

**FOR FURTHER INFORMATION CONTACT:** Denver Presley, Jr., Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., P150–400B, Rockville, MD 20850, 301–796–3793.

**SUPPLEMENTARY INFORMATION:** On August 23, 2011, the Agency submitted a proposed collection of information entitled “Temporary Marketing Permit Applications” to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0133. The approval expires on September 30, 2014. A copy of the supporting statement for this information collection is available on the Internet at http://www.reginfo.gov/public/do/PRAMain.