EXEMPTIONS CLAIMED FOR SYSTEM:

Pursuant to 5 U.S.C. 552a(k)(2), records in this system are exempt from the requirements of subsections (c)(3); (d); (e)(1); (e)(4)(G), (H), (I); and (f) of 5 U.S.C. 552a, provided, however, that if any individual is denied any right, privilege, or benefit that he or she would otherwise be entitled to by Federal law, or for which he or she would otherwise be eligible, as a result of the maintenance of these records, such material shall be provided to the individual, except to the extent that the disclosure of the material would reveal the identity of a source who furnished information to the Government with an express promise that the identity of the source would be held in confidence.

[FR Doc. 2015–17924 Filed 7–21–15; 8:45 am]

BILLING CODE 6760–01–P

GENERAL SERVICES ADMINISTRATION

[Notice-CECANF–2015–07; Docket No. 2015–0004; Sequence No. 7]

Commission To Eliminate Child Abuse and Neglect Fatalities; Announcement of Meetings

AGENCY: Commission to Eliminate Child Abuse and Neglect Fatalities.

ACTION: Meeting Notice.

SUMMARY: The Commission to Eliminate Child Abuse and Neglect Fatalities (CECANF), a Federal Advisory Committee established by the Protect Our Kids Act of 2012, Public Law 112–275, will hold a meeting open to the public on Thursday, August 6, 2015 and Friday, August 7, 2015 in New York, New York.

DATES: The meeting will be held on Thursday, August 6, 2015, from 8:00 a.m. to 5:30 p.m., Eastern Daylight Time, and Friday, August 7, 2015, from 8:00 a.m. to 12:30 p.m., Eastern Daylight Time.

ADDRESSES: CECANF will convene its meeting at the ACS Children's Center Auditorium, 492 First Avenue at 28th Street, New York, NY 10016. This site is accessible to individuals with disabilities. The meeting also will be made available via teleconference and/or webinar.

Submit comments identified by ''Notice-CECANF–2015–07,'' by either of the following methods:

• Regulations.gov: *http://www.regulations.gov.*

Submit comments via the Federal eRulemaking portal by searching for ''Notice-CECANF–2015–07.'' Select the link ''Comment Now'' that corresponds with ''Notice-CECANF–2015–07.'' Follow the instructions provided on the screen. Please include your name, organization name (if any), and ''Notice-CECANF–2015–07'' on your attached document.

• *Mail:* U.S. General Services Administration, 1800 F Street NW., Room 7003D, Washington DC 20405, Attention: Tom Hodnett (CD) for CECANF.

Instructions: Please submit comments only and cite ''Notice-CECANF–2015–07'' in all correspondence related to this notice. All comments received will be posted without change to *http://www.regulations.gov,* including any personal and/or business confidential information provided.

FOR FURTHER INFORMATION CONTACT: Visit the CECANF Web site at *https://eliminatechildabusefatalities.sites.usa.gov/* or contact Patricia Brincefield, Communications Director, at 202–818–9596, U.S. General Services Administration, 1800 F Street NW., Room 7003D, Washington DC 20405, Attention: Tom Hodnett (CD) for CECANF.

SUPPLEMENTARY INFORMATION:

Background: CECANF was established to develop a national strategy and recommendations for reducing fatalities resulting from child abuse and neglect.

Agenda: This meeting will explore key research, policy, and practice in New York City related to addressing and preventing child abuse and neglect fatalities. Commission members will then continue discussing the work plans of the Commission subcommittees, the information that they have obtained to date, and emerging recommendations.

Attendance at the Meeting: Individuals interested in attending the meeting in person or participating by webinar and teleconference must register in advance. To register to attend in person or by webinar/phone, please go to *http://meetingtomorrow.com/webcast/CECANFNY* and follow the prompts. Once you register, you will receive a confirmation email with the webinar login and teleconference number. Detailed meeting minutes will be posted within 90 days of the meeting. Members of the public will not have the opportunity to ask questions or otherwise participate in the meeting.

However, members of the public wishing to comment should follow the steps detailed under the heading **ADDRESSES** in this publication or contact us via the CECANF Web site at *https://eliminatechildabusefatalities.sites.usa.gov/contact-us/.*

Dated: July 15, 2015.

Karen White,

Executive Assistant.

[FR Doc. 2015–17954 Filed 7–21–15; 8:45 am]

BILLING CODE 6820–34–P

GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–0262; Docket 2015–0001; Sequence 9]

General Services Administration Acquisition Regulation; Submission for OMB Review; Identification of Products With Environmental Attributes

AGENCY: Office of Acquisition Policy, General Services Administration (GSA).

ACTION: Notice of request for comments regarding an extension of a previously existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act the Regulatory Secretariat Division 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 identification of products with environmental attributes. A notice was published in the **Federal Register** at 80 FR 22351 on April 28, 2015. No comments were received.

DATES: Submit comments on or before: August 21, 2015.

ADDRESSES: Submit comments identified by Information Collection 3090–0262, Identification of Products with Environmental Attributes, by any of the following methods:

• *Regulations.gov: http://www.regulations.gov.* Submit comments via the Federal eRulemaking portal by inputting ''Information Collection 3090–0262, Identification of Products with Environmental Attributes'', under the heading ''Enter Keyword or ID'' and selecting ''Search''. Select the link ''Submit a Comment'' that corresponds with ''Information Collection 3090–0262, Identification of Products with Environmental Attributes''. Follow the instructions provided at the ''Submit a Comment'' screen. Please include your name, company name (if any), and ''Information Collection 3090–0262, Identification of Products with Environmental Attributes'' on your attached document.

• *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street, NW., Washington, DC 20405. ATTN: Ms. Flowers/IC 3090–0262, Identification of

Products with Environmental Attributes.

Instructions: Please submit comments only and cite Information Collection 3090–0262, Identification of Products with Environmental Attributes, 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.

FOR FURTHER INFORMATION CONTACT: Ms. Dana Munson, Procurement Analyst, General Services Acquisition Policy Division, GSA, at telephone 202–357–9652 or via email to *dana.munson@gsa.gov*.

SUPPLEMENTARY INFORMATION:

A. Purpose

The General Services Administration (GSA) requires contractors holding Multiple Award Schedule Contracts to identify in their GSA price lists those products that they market commercially that have environmental attributes in accordance with GSAR clause 552.238–72. The identification of these products will enable Federal agencies to maximize the use of these products and meet the responsibilities expressed in statutes and executive orders.

B. Annual Reporting Burden

Respondents: 9,000.
Responses per Respondent: 1.
Annual Responses: 9,000.
Hours per Response: 1.
Total Burden Hours: 9,000.

C. Public Comments

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.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 3090–0262, Identification of Products with Environmental Attributes, in all correspondence.

Dated: July 16, 2015.

Jeffrey A. Koses,

Senior Procurement Executive, Director, Office of Acquisition Policy.

[FR Doc. 2015–17904 Filed 7–21–15; 8:45 am]

BILLING CODE 6820–61–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–0736]

Obstetrics and Gynecology Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committee: Obstetrics and Gynecology Devices Panel of the Medical Devices Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the Agency on FDA's regulatory issues.

Date and Time: The meeting will be held on September 24, 2015, from 8 a.m. to 6 p.m.

ADDRESSES: FDA is opening a docket for interested persons to submit electronic or written comments regarding this meeting. The docket number is FDA–2014–N–0736. Please see the *Procedure* section of the notice for further information.

Location: FDA White Oak Campus, 10903 New Hampshire Ave., Bldg. 31 Conference Center, the Great Room (Rm. 1503), Silver Spring, MD 20993–0002. Answers to commonly asked questions including information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: *http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm.*

Contact Person: Shanika Craig, Center for Devices and Radiological Health (CDRH), Food and Drug Administration, 10903 New Hampshire Ave., Silver Spring, MD 20993, 301–796–6639, *Shanika.Craig@fda.hhs.gov,* or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area). A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's Web site at *http://www.fda.gov/AdvisoryCommittees/default.htm* and scroll down to the appropriate advisory committee meeting link, or call the advisory committee information line to learn about possible modifications before coming to the meeting.

Agenda: On September 24, 2015, the committee will discuss the risks and benefits of Bayer HealthCare's Essure System for permanent female sterilization. The system, originally approved in November 2002, under P020014, consists of a delivery system and nickel-containing permanent implants. The implants are placed without a skin incision, through the vagina, within each fallopian tube; they elicit tissue ingrowth, which over time results in tubal occlusion.

FDA is convening this committee to seek expert scientific and clinical opinion on the risks and benefits of the Essure System. The committee will be asked to evaluate currently available scientific data pertaining to the safety and effectiveness of the Essure System, such as events related to implant perforation/migration, device removal, chronic pain, allergic reactions, and unintended pregnancy. The committee will be asked to provide recommendations regarding appropriate device use, product labeling, and potential need for additional postmarket clinical studies.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at *http://www.fda.gov/AdvisoryCommittees/Calendar/default.htm.* Scroll down to the appropriate advisory committee meeting link.

CDRH plans to provide a live Webcast of the September 24, 2015, meeting of the Obstetrics and Gynecology Devices Panel. While CDRH is working to make Webcasts available to the public for all advisory committee meetings held at the White Oak campus, there are instances where the Webcast transmission is not successful; staff will work to re-establish the transmission as soon as possible. The link for the Webcast is available at: *https://collaboration.fda.gov/gudpm052015/.* Further information regarding the Webcast, including the Web address for the Webcast, will be made available at least 2 days in advance of the meeting at the following Web site: *https://collaboration.fda.gov/ogdp2015/.*

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending