institutions to have written policies and procedures in place to ensure that their employees who perform mortgage loan originations comply with the registration and other SAFE Act requirements.

There is no change in the method or substance of the collection. The overall reduction in burden hours is the result of economic fluctuation. In particular, the number of respondents has decreased while the hours per response and frequency of responses have remained the same.

Dated at Washington, DC, on March 4, 2019.

Federal Deposit Insurance Corporation.

Robert E. Feldman,

Executive Secretary.

[FR Doc. 2019–04139 Filed 3–6–19; 8:45 am]

BILLING CODE 6714–01–P

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0073; Docket No. 2019–0003; Sequence No. 15]

Information Collection; Advance Payments

AGENCY: Department of Defense (DOD), General Services Administration (GSA), and National Aeronautics and Space Administration (NASA).

ACTION: Notice of request for an extension of an information collection requirement for an 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 advanced payments.

DATES: *Submit comments on or before:* May 6, 2019.

ADDRESSES: The FAR Council invites interested persons to submit comments on this collection by any of the following methods:

- *Federal eRulemaking Portal:* This website provides the ability to type short comments directly into the comment field or attach a file for lengthier comments. Go to *http://www.regulations.gov* and follow the instructions on the site.
- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW, Washington, DC 20405. ATTN: Ms. Mandell/IC 9000–0073, Advance Payments.

Instructions: Please submit comments only and cite Information Collection 9000–0073, Advance Payments, in all correspondence related to this collection. Comments received generally will be posted without change to *http://www.regulations.gov,* including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check *www.regulations.gov,* approximately two to three days after submission to verify posting (except allow 30 days for posting of comments submitted by mail).

FOR FURTHER INFORMATION CONTACT: Mr. Kevin Funk, Procurement Analyst, at telephone 202–357–5805, or via email at *kevin.funk@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

Advance payments may be authorized under Federal contracts and subcontracts. Advance payments are the least preferred method of contract financing and require special determinations by the agency head or designee. Specific financial information about the contractor is required before such payments can be authorized (see FAR 32.4 and 52.232–12). The information is used to determine if advance payments should be provided to the contractor.

B. Annual Reporting Burden

Respondents: 73.

Responses per Respondent: 12.

Annual Responses: 876.

Hours per Response: 1.42.

Total Burden Hours: 1,244.

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 General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW, Washington, DC 20405, at 202–501–4755. Please cite OMB Control No. 9000–0073, Advance Payments, in all correspondence.

Janet Fry,

Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2019–04100 Filed 3–6–19; 8:45 am]

BILLING CODE 6820–EP–P

GENERAL SERVICES ADMINISTRATION

[Notice–WSCC–2019–02; Docket 2019–0004; Sequence No. 2]

Women's Suffrage Centennial Commission; Notification of Public Meetings

AGENCY: Women's Suffrage Centennial Commission, General Services Administration.

ACTION: Meetings notice.

SUMMARY: Notice of two meetings is being provided according to the requirements of the Federal Advisory Committee Act. This notice provides the schedule and agenda for the March 27, 2019, and June 3, 2019, meetings of the Women's Suffrage Centennial Commission (Commission). The meetings are open to the public.

DATES: *Meeting dates:* The first meeting will be held on Wednesday, March 27, 2019, beginning at 1:00 p.m., and ending no later than 3:00 p.m. (Eastern Daylight Time). The second meeting will be held on Monday, June 3, 2019, beginning at 9:00 a.m., and ending no later than 4:00 p.m. (Eastern Daylight Time).

ADDRESSES: The meeting on March 27, 2019, will be a telephonic meeting. The public may dial into the meeting by calling 866–660–8781 participant code: 9420503. The meeting on June 3, 2019, will be held at the Department of the Interior, 1849 C Street NW, Washington, DC 20240, in the South Penthouse on the 7th Floor.

FOR FURTHER INFORMATION CONTACT: Kim Oliver, Designated Federal Officer, Women's Suffrage Centennial Commission, 1849 C Street NW, Room 7313, Washington, DC 20240; phone: 202–912–7510; fax: 202–219–2100; email: *kmoliver@blm.gov.*

SUPPLEMENTARY INFORMATION:

Background

Congress passed legislation to create the Women's Suffrage Centennial Commission Act, a bill, ''to ensure a suitable observance of the centennial of the passage and ratification of the 19th

Amendment of the Constitution of the United States providing for women's suffrage.''

The duties of the Commission, as written in the law, include: (1) To encourage, plan, develop, and execute programs, projects, and activities to commemorate the centennial of the passage and ratification of the 19th Amendment; (2) To encourage private organizations and State and local Governments to organize and participate in activities commemorating the centennial of the passage and ratification of the 19th Amendment; (3) To facilitate and coordinate activities throughout the United States relating to the centennial of the passage and ratification of the 19th Amendment; (4) To serve as a clearinghouse for the collection and dissemination of information about events and plans for the centennial of the passage and ratification of the 19th Amendment; and (5) To develop recommendations for Congress and the President for commemorating the centennial of the passage and ratification of the 19th Amendment.

Meeting Agenda for March 27, 2019

- Welcome and Introductions
- Executive Director update
- Subcommittee updates
- Public Comment Period
- Adjourn

Meeting Agenda for June 3, 2019

- Welcome and Introductions
- Commission business and administrative items
- Executive Director update
- Presentations from informative speakers
- Subcommittee updates
- Public Comment Period
- 2019 Meeting Schedule
- Adjourn

The meetings are open to the public, but preregistration is required. Any individual who wishes to attend the meeting should register via email at *kmoliver@blm.gov* or telephone 202–912–7510. Space is limited and requests to attend will be accommodated in the order they are received.

Interested persons may choose to make a public comment at the meeting during the designated time for this purpose. Public comments shall be limited by minutes based on the number of participants signed up to comment for the allotted time, and subject to agenda time changes based on the speed of the commission's work through the agenda. Speakers who wish to expand upon their oral statements, or those who had wished to speak but could not be accommodated on the agenda, may submit written statements up to 30 days after the meeting.

Members of the public may also choose to submit written comments by mailing them to Kim Oliver, Designated Federal Officer, 1849 C Street NW, Room 7313, Washington, DC 20240, or via email at *kmoliver@blm.gov.* Please contact Ms. Oliver at the email address above to obtain meeting materials. All written comments received will be provided to the Commission. Detailed minutes of the meeting will be available for public inspection within 90 days of the meeting.

Individuals requiring special accommodations to access the public meeting should contact Ms. Oliver at least five business days prior to each meeting, so that appropriate arrangements can be made.

Public Disclosure of Comments

Before including your address, phone number, email address, or other personal identifying information in your comment, you should be aware that your entire comment—including your personal identifying information—may be made publicly available at any time. While you can ask us in your comment to withhold your personal identifying information from public review, we cannot guarantee that we will be able to do so.

Jeffrey A. Koses,

Senior Procurement Executive, Office of Acquisition Policy, Office of Governmentwide Policy, General Services Administration.

[FR Doc. 2019–04143 Filed 3–6–19; 8:45 am]

BILLING CODE 3420–37–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Nos. FDA–2017–M–6870, FDA–2018–M–3584, and FDA–2018–M–3870]

Medical Devices Regulated by the Center for Biologics Evaluation and Research; Availability of Safety and Effectiveness Summaries for Premarket Approval Applications

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is publishing a list of premarket approval applications (PMAs) that have been approved by the Center for Biologics Evaluation and Research (CBER). This list is intended to inform the public of the availability of safety and effectiveness summaries of approved PMAs through the internet and FDA's Dockets Management Staff.

ADDRESSES: You may submit either electronic or written comments at any time as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Since your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov.*
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand Delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment as well as any attachments, except for information submitted, marked, and identified as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket Nos. FDA–2017–M–6870, FDA–2018–M–3584, and FDA–2018–M–3870 for ''Medical Devices Regulated by the Center for Biologics Evaluation and Research; Availability of Safety and Effectiveness Summaries for Premarket Approval Applications.'' Received comments will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9