and Receiverships, Attention: Receivership Oversight Department 34.6, 1601 Bryan Street, Dallas, TX 75201.

No comments concerning the termination of this receivership will be considered which are not sent within this time frame.

Dated: August 23, 2017.

Federal Deposit Insurance Corporation.

Robert E. Feldman,

Executive Secretary.

[FR Doc. 2017–18156 Filed 8–25–17; 8:45 am]

BILLING CODE 6714–01–P

FEDERAL DEPOSIT INSURANCE CORPORATION

Notice to All Interested Parties of the Termination of the Receivership of 10301—First Suburban National Bank, Maywood, Illinois

Notice is hereby given that the Federal Deposit Insurance Corporation (FDIC) as Receiver for First Suburban National Bank, Maywood, Illinois (''the Receiver'') intends to terminate its receivership for said institution. The FDIC was appointed Receiver of First Suburban National Bank on October 22, 2010. The liquidation of the receivership assets has been completed. To the extent permitted by available funds and in accordance with law, the Receiver will be making a final dividend payment to proven creditors.

Based upon the foregoing, the Receiver has determined that the continued existence of the receivership will serve no useful purpose. Consequently, notice is given that the receivership shall be terminated, to be effective no sooner than thirty days after the date of this notice. If any person wishes to comment concerning the termination of the receivership, such comment must be made in writing and sent within thirty days of the date of this notice to: Federal Deposit Insurance Corporation, Division of Resolutions and Receiverships, Attention: Receivership Oversight Department 34.6, 1601 Bryan Street, Dallas, TX 75201.

No comments concerning the termination of this receivership will be considered which are not sent within this time frame.

Dated August 22, 2017.

Federal Deposit Insurance Corporation.

Robert E. Feldman,

Executive Secretary.

[FR Doc. 2017–18109 Filed 8–25–17; 8:45 am]

BILLING CODE 6714–01–P

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0180; Docket No. 2017–0053; Sequence 12]

Information Collection; Affirmative Procurement of Biobased Procurements Under Services and Construction Contracts

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

ACTION: Notice of request for public comments regarding an extension to an existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat Division (MVCB) 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 Biobased Procurements.

DATES: Submit comments on or before October 27, 2017.

ADDRESSES: Submit comments identified by Information Collection 9000–0180, Affirmative Procurement of Biobased Procurements Under Services and Construction Contracts, by any of the following methods:

- *Regulations.gov: http://www.regulations.gov.* Submit comments via the Federal eRulemaking portal by searching the OMB control number 9000–0180. Select the link ''Comment Now'' that corresponds with ''Information Collection 9000–0180, Affirmative Procurement of Biobased Procurements Under Services and Construction Contracts. Follow the instructions provided on the screen. Please include your name, company name (if any), and ''Information Collection 9000–0180, ''Affirmative Procurement of Biobased Procurements Under Services and Construction Contracts'' on your attached document.
- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Sosa/IC 9000–0180, Biobased Procurements.

Instructions: Please submit comments only and cite Information Collection 9000–0180, Affirmative Procurement of Biobased Procurements Under Services and Construction Contracts. 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. Charles Gray, Procurement Analyst, Office of Governmentwide Acquisition Policy, at telephone 703–795–6328, or email *charles.gray@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

Federal Acquisition Regulation clause 52.223–2, Affirmative Procurement of Biobased Products Under Service and Construction Contracts, requires prime contractors to report annually the product types and dollar values of U.S. Department of Agriculture (USDA)-designated biobased products purchased to the System for Award Management (SAM) Web site. The information reported by prime contractors enables Federal agencies to report annually to the Office of Federal Procurement Policy (OFPP) concerning actions taken to implement and measure progress in carrying out the preference for biobased products required under section 9002 of the Farm Security and Rural Investment Act of 2002, codified at 7 U.S.C. 8102.

B. Annual Reporting Burden

To determine the number of contractors performing construction and service contracts that may involve the purchase of USDA-designated biobased products, fiscal year 2016 data in the Federal Procurement Data System (FPDS) was reviewed to calculate the number entities with unique DUNS numbers that were awarded contracts for the following selected Product Services Codes: A—Research and Development; F—Natural Resources Management; J—Maintenance, Repair, and Rebuilding of Equipment; M—Operation of Government-Owned Facility; S—Utilities and Housekeeping Services; T—Photographic, Mapping, Printing, and Publication Services; Y—Construction of Structures and Facilities; and Z—Maintenance, Repair or Alteration of Real Property. The clause at FAR 52.223–2 will apply to the majority of the contract actions in the selected PSCs

The estimated total burden is as follows:

Respondents: 51,457.

Responses per Respondent: 5.

Total Annual Responses: 257,285.

Hours per Response: 5.

Total Burden Hours: 1,286,425.

Affected Public: Businesses or other for-profit and not-for-profit institutions.

Frequency: Annually.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary for the proper performance of functions of the FAR, 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; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

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, telephone 202–501–4755. Please cite OMB Control No. 9000–0180, Affirmative Procurement of Biobased Procurements Under Services and Construction Contracts, in all correspondence.

Dated: August 22, 2017.

Lorin S. Curit,

Director, Federal Acquisition Policy Division, Office of Government-wide Acquisition Policy, Office of Acquisition Policy, Office of Government-wide Policy.

[FR Doc. 2017–18105 Filed 8–25–17; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2017–N–0001]

Cellular, Tissue, and Gene Therapies Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) announces a forthcoming public advisory committee meeting of the Cellular, Tissue, and Gene Therapies Advisory Committee (CTGTAC). The general function of the committee is to provide advice and recommendations to the Agency on FDA's regulatory issues. The meeting will be open to the public.

DATES: The meeting will be held on October 12, 2017, from 8:30 a.m. to 5 p.m.

ADDRESSES: FDA White Oak Campus, 10903 New Hampshire Ave., Bldg. 31 Conference Center, the Great Room (Rm. 1503) Silver Spring, MD 20993–0002.

For those unable to attend in person, the meeting will also be Webcast and will be available at the following link: *https://collaboration.fda.gov/ctgtac101217.* Answers to commonly asked questions including information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: *https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm.*

FOR FURTHER INFORMATION CONTACT: Prabhakara L. Atreya or Denise Royster, Center for Biologics Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 6306, Silver Spring, MD 20993–0002, 240–402–8006, *prabhakara.atreya@fda.hhs.gov* and 240–402–8158, *denise.royster@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 *https://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.

SUPPLEMENTARY INFORMATION:

Agenda: On October 12, 2017, the CTGTAC will meet in an open session to discuss and make recommendations on the safety and effectiveness of biologics license application (BLA) for voretigene neparvovec (BLA 125610), submitted by Spark Therapeutics, Inc. The proposed indication (use) for this product is for the treatment of patients with vision loss due to confirmed biallelic RPE65 mutation-associated retinal dystrophy.

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 *https://www.fda.gov/AdvisoryCommittees/Calendar/default.htm.* Scroll down to the appropriate advisory committee meeting link.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before October 4, 2017. Oral presentations from the public will be scheduled between approximately 11:15 a.m. and 12:15 p.m. Those individuals interested in making formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before September 26, 2017. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by September 27, 2017.

Persons attending FDA's advisory committee meetings are advised that the Agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with disabilities. If you require accommodations due to a disability, please contact Prabhakara Atreya at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at: *https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm* for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).