

CONTACT PERSON FOR MORE INFORMATION:
David Eng, Secretary; Phone: (202) 523–5725; Email: secretary@fmc.gov.

David Eng,
Secretary.

[FR Doc. 2026–21259 Filed 10–15–26; 11:15 am]

BILLING CODE 6730–02–P

FEDERAL RETIREMENT THRIFT INVESTMENT BOARD

Notice of Board Meeting

DATES: October 27, 2026 at 9:00 a.m. ET.

ADDRESSES: Telephonic. Dial-in (listen only) information: Number: 1–202–599–1426, Code: 192 871 229#; or via web: <https://www.frtib.gov/>.

FOR FURTHER INFORMATION CONTACT:

James Kaplan, Director, Office of External Affairs, (202) 864–7150.

SUPPLEMENTARY INFORMATION:

Board Meeting Agenda

Open Session

1. Approval of the September 22, 2026, Board Meeting Minutes
2. Monthly Reports
 - (a) Participant Report
 - (b) Legislative Report
3. Quarterly Reports
 - (c) Investment Review
 - (d) Audit Status
 - (e) Budget Review
4. Internal Audit Update
5. Participant Satisfaction Survey Report
6. Social Science Update
7. OEA Office Presentation

Closed Session

8. Information Covered under 5 U.S.C. 552b (c)(6), (c)(9)(B), and (c)(10).
Authority: 5 U.S.C. 552b (e)(1).
Dated: October 15, 2026.

Dharmesh Vashee,

General Counsel, Federal Retirement Thrift Investment Board.

[FR Doc. 2026–21269 Filed 10–16–26; 8:45 am]

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GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–0322; Docket No. 2026–0564; Sequence No. 1]

Information Collection; General Services Administration Acquisition Regulation; Prohibition on Certain Supply Chain Services or Equipment Under Lease Acquisitions and Commercial Solution Openings

AGENCY: Office of the Chief Acquisition Officer, General Services Administration (GSA).

ACTION: Notice; request for comments.

SUMMARY: In accordance with the Paperwork Reduction Act of 1995 and Office of Management and Budget (OMB) regulations, GSA is extending a previously approved information collection regarding OMB Control Number 3090–0322, Prohibition on Certain Supply Chain Services or Equipment Under Lease Acquisitions and Commercial Solution Openings, without changes except for updated wage rates, until the information collection can be assessed and revised after the completion of the rulemaking related to the Revolutionary GSA Regulation (GSAR) Overhaul (RGO).

DATES: GSA will consider all comments received by December 18, 2026.

ADDRESSES: GSA invites interested persons to submit comments on this collection through <https://www.regulations.gov> and follow the instructions on the site. This website provides the ability to type short comments directly into the comment field or attach a file for lengthier comments. If there are difficulties submitting comments, contact the GSA Regulatory Secretariat Division at GSARegSec@gsa.gov.

Instructions: All items submitted must cite OMB Control No. 3090–0322, Prohibition on Certain Supply Chain Services or Equipment Under Lease Acquisitions and Commercial Solution Openings. Comments received generally will be posted without change to <https://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.

FOR FURTHER INFORMATION CONTACT:

Amy Lara, Procurement Analyst, General Services Acquisition Policy Division, at telephone 816–589–3783, or via email at gsarpolicy@gsa.gov.

SUPPLEMENTARY INFORMATION:

A. OMB Control Number, Title, and Any Associated Form(s)

OMB Control No. 3090–0322, Prohibition on Certain Supply Chain Services or Equipment Under Lease Acquisitions and Commercial Solution Openings.

B. Needs and Uses

There are two purposes. The first (“889”) supports implementation of Section 889 of the John S. McCain National Defense Authorization Act for Fiscal Year 2019 (Pub. L. 115–232) under lease acquisitions and

commercial solution openings. This section prohibits agencies from procuring, obtaining, extending or renewing a contract with contractors that will provide or use covered telecommunication equipment or services as a substantial or essential component of any system, or as a critical technology as part of any system on or after August 13, 2020, unless an exception applies.

The second (“FASCSA Orders”) supports implementation of supply chain risk information sharing and exclusion or removal orders consistent with the Federal Acquisition Supply Chain Security Act of 2018 and a final rule issued by the Federal Acquisition Security Council. The implementation of supply chain risk information sharing and exclusion or removal orders FAR interim rule requires complying with exclusion or removal orders (“FASCSA Orders”) and sharing certain supply chain risk information with the Federal Acquisition Security Council (FASC) when applicable FASCSA orders are issued from one or a combination of the following FASCSA orders-issuing agencies: Department of Homeland Security (DHS), the Department of Defense (DoD), and/or the Office of the Director of National Intelligence (DNI). Only DHS may issue orders applicable to GSA (*i.e.*, civilian agencies).

For 889, the requirement is implemented in the Federal Acquisition Regulation (FAR) through the provision at FAR 52.204–24, Representation Regarding Certain Telecommunications and Video Surveillance Services or Equipment and the clause at FAR 52.204–25, Prohibition on Contracting for Certain Telecommunications and Video Surveillance Services or Equipment.

For FASCSA Orders, the requirement is implemented in the FAR through the provision at FAR 52.204–29, Federal Acquisition Supply Chain Security Act Orders—Representation and Disclosures and the clause at FAR 52.204–30, Federal Acquisition Supply Chain Security Act Orders—Prohibition.

C. Annual Burden

1. FAR 52.204–24 for GSA Lease Acquisitions

Respondents: 3,100.
Responses per Respondent: 1.
Total Responses: 3,100.
Hours per Response: 1.5.
Total Burden Hours: 4,650.

2. FAR 52.204–25 for GSA Lease Acquisitions

Respondents: 62.
Responses per Respondent: 1.

Total Responses: 62.
Hours per Response: 1.5.
Total Burden Hours: 93.

3. FAR 52.204–29 for GSA Lease Acquisitions

Respondents: 186.
Responses per Respondent: 1.
Total Responses: 186.
Hours per Response: 2.
Total Burden Hours: 372.

4. FAR 52.204–30 for GSA Lease Acquisitions

Respondents: 124.
Responses per Respondent: 1.
Total Responses: 124.
Hours per Response: 2.
Total Burden Hours: 248.

Note: GSA solicits and awards so few CSO procurements (on average less than 5 per year), the burden is negligible and therefore not included in this estimate. *Obtaining Copies:* Requesters may obtain a copy of the information collection documents from the GSA Regulatory Secretariat Division, by emailing GSARegSec@gsa.gov. Please cite “Information Collection 3090–0322”, in all correspondence.

Richard Speidel,
Deputy Chief Data Officer, General Services Administration.

[FR Doc. 2026–21255 Filed 10–16–26; 8:45 am]

BILLING CODE:P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1784]

Bulk Manufacturer of Controlled Substances Application: Scottsdale Research Institute

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Scottsdale Research Institute has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to Supplementary Information listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants, therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before December 18, 2026. Such persons may also file a written request for a hearing on the application on or before December 18, 2026.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short

comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to <https://www.regulations.gov> and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on <https://www.regulations.gov>. If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on September 8, 2026, Scottsdale Research Institute, 12815 North Cave Creek Road, Phoenix, Arizona 85022, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
lbgaine	7260	I

The company plans to bulk manufacture the listed controlled substance for internal research purposes and to support clinical trials. No other activity for this drug code is authorized for this registration.

Justin Wood,
Deputy Assistant Administrator.

[FR Doc. 2026–21360 Filed 10–16–26; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1788]

Bulk Manufacturer of Controlled Substances Application: Janssen Pharmaceuticals, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Janssen Pharmaceuticals, Inc. has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to

SUPPLEMENTARY INFORMATION listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants, therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before December 18, 2026. Such persons may also file a written request for a hearing on the application on or before December 18, 2026.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to <https://www.regulations.gov> and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on <https://www.regulations.gov>. If you have received a Comment Tracking Number, your comment has been

successfully submitted and there is no need to resubmit the same comment.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on September 9, 2026, Janssen Pharmaceuticals, Inc., 1440 Olympic Drive, Buildings 1 and 5 and 8 and 20, Athens, Georgia 30601–1645, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Methylphenidate	1724	II

The company plans to manufacture the listed controlled substance for the internal use intermediates for sale to its customers. No other activity for this drug code is authorized for this registration.

Justin Wood,
Deputy Assistant Administrator.

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