Register pursuant to Section 6(b) of the Act on November 12, 2019 (84 FR 61071).

The last notification was filed with the Department on July 7, 2020. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on July 31, 2020 (85 FR 46179).

Suzanne Morris,

Chief, Premerger and Division Statistics, Antitrust Division.

[FR Doc. 2020–25745 Filed 11–20–20; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–745]

Bulk Manufacturer of Controlled Substances Application: Johnson Matthey Pharmaceutical Materials Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Johnson Matthey Pharmaceutical Materials Inc. has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTAL INFORMATION** listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before January 22, 2021. Such persons may also file a written request for a hearing on the application on or before January 22, 2021.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on November 2, 2020, Johnson Matthey Pharmaceutical Materials Inc., 25 Patton Road, Devens, Massachusetts 01434, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Amphetamine	1100	II
Methylphenidate	1724	II
Nabilone	7379	II
Hydrocodone	9193	II
Levorphanol	9220	II
Thebaine	9333	II
Alfentanil	9737	II
Remifentanil	9739	II
Sufentanil	9740	II

The company plans to support its other manufacturing facilities located in West Deptford, New Jersey and Conshohocken, Pennsylvania with manufacturing and analytical testing.

In reference to drug code 9333 as bulk, the company plans to manufacture a Thebaine derivative for distribution to its customers. No other activity for these drug codes is authorized for this registration.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–25764 Filed 11–20–20; 8:45 am]

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OCCUPATIONAL SAFETY AND HEALTH ADMINISTRATION

[Docket No. OSHA–2011–0028]

Grain Handling Facilities; Extension of the Office of Management and Budget's (OMB) Approval of Information Collection (Paperwork) Requirements

AGENCY: Occupational Safety and Health Administration (OSHA), Labor.

ACTION: Request for public comments.

SUMMARY: OSHA solicits public comments concerning the proposal to extend the Office of Management and Budget's (OMB) approval of the information collection requirements contained in the standard on Grain Handling Facilities.

DATES: Comments must be submitted (postmarked, sent, or received) by January 22, 2021.

ADDRESSES:

Electronically: You may submit comments and attachments electronically at *http://www.regulations,gov,* which is the Federal eRulemaking Portal. Follow the instructions online for submitting comments.

Facsimile: If your comments, including attachments, are not longer than 10 pages you may fax them to the OSHA Docket Office at (202) 693–1648.

Mail, hand delivery, express mail, messenger, or courier service: When using this method, you must submit a copy of your comments and attachments to the OSHA Docket Office, OSHA Docket No. OSHA–2011–0028, Occupational Safety and Health Administration, U.S. Department of Labor, Room N–3653, 200 Constitution Avenue NW, Washington, DC 20210. *Please note:* While OSHA's Docket Office is continuing to accept and process submissions by regular mail, due to the COVID–19 pandemic, the Docket Office is closed to the public and not able to received submissions to the docket by hand, express mail, messenger, and courier service.

Instructions: All submissions must include the agency name and the OSHA docket number (OSHA–2011–0028) for the Information Collection Request (ICR). All comments, including any personal information you provide, such as social security numbers and date of birth, are placed in the public docket without change, and may be made available online at *http://www.regulations.gov*. For further information on submitting comments see the ''Public Participation'' heading in the section of this notice titled **SUPPLEMENTARY INFORMATION**.

Docket: To read or download comments or other material in the docket, go to *http://www.regulations.gov* or the OSHA Docket Office at the above address. All documents in the docket (including this **Federal Register** notice) are listed in the *http://www.regulations.gov* index; however, some information (*e.g.,* copyrighted material) is not publicly available to read or download through the website. All submissions, including copyrighted material, are available for inspection and copying at the OSHA Docket Office. You may also contact Theda Kenney at the below phone number to obtain a copy of the ICR.

FOR FURTHER INFORMATION CONTACT: Seleda Perryman or Theda Kenney, Directorate of Standards and Guidance, OSHA, U.S. Department of Labor; telephone (202) 693–2222.

SUPPLEMENTARY INFORMATION:

I. Background

The Department of Labor, as part of the continuing effort to reduce paperwork and respondent (*i.e.,* employer) burden, conducts a preclearance consultation program to provide the public with an opportunity to comment on proposed and continuing information collection requirements in accordance with the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3506(c)(2)(A)). This program ensures that information is in the desired format, reporting burden (time and costs) is minimal, collection instruments are clearly understood, and OSHA's estimate of the information collection burden is accurate. The Occupational Safety and Health Act of 1970 (OSH Act) (29 U.S.C. 651 *et seq.*)