

exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Office of Diversion Control (“Assistant Administrator”) pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on September 5, 2016, Johnson Matthey Inc., Pharmaceutical Materials, 2003 Nolte Drive, West Deptford, New Jersey 08066-1742, applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Coca Leaves .....	9040	II
Thebaine .....	9333	II
Opium, raw .....	9600	II
Noroxymorphone .....	9668	II
Poppy Straw Concentrate .....	9670	II
Fentanyl .....	9801	II

The company plans to import Thebaine derivatives and Fentanyl as reference standards. The company plans to import the remaining listed controlled substances as raw materials, to be used in the manufacture of bulk controlled substances, for distribution to its customers.

Dated: October 11, 2016.

**Louis J. Milione,**  
Assistant Administrator.

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Bulk Manufacturer of Controlled Substances Application: Johnson Matthey Inc.**

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.33(a) on or before December 19, 2016.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement

Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** The Attorney General has delegated her authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Office of Diversion Control (“Assistant Administrator”) pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.33(a), this is notice that on September 5, 2016, Johnson Matthey Inc., Custom Pharmaceuticals Department, 2003 Nolte Drive, West Deptford, New Jersey 08066-1742 applied to be registered as a bulk manufacturer the following basic classes of controlled substances:

Controlled Substance	Drug code	Schedule
Gamma Hydroxybutyric Acid .....	2010	I
Marihuana .....	7360	I
Tetrahydrocannabinols .....	7370	I
Dihydromorphine .....	9145	I
Difenoxin .....	9168	I
Propiram .....	9649	I
Amphetamine .....	1100	II
Methamphetamine .....	1105	II
Lisdexamfetamine .....	1205	II
Methylphenidate .....	1724	II
Nabilone .....	7379	II
Cocaine .....	9041	II
Codeine .....	9050	II
Dihydrocodeine .....	9120	II
Oxycodone .....	9143	II
Hydromorphone .....	9150	II
Diphenoxylate .....	9170	II
Ecgonine .....	9180	II
Hydrocodone .....	9193	II
Meperidine .....	9230	II
Methadone .....	9250	II
Methadone intermediate .....	9254	II
Morphine .....	9300	II
Thebaine .....	9333	II
Oxymorphone .....	9652	II
Noroxymorphone .....	9668	II
Alfentanil .....	9737	II
Remifentanil .....	9739	II
Sufentanil .....	9740	II
Tapentadol .....	9780	II
Fentanyl .....	9801	II

The company plans to manufacture the listed controlled substances in bulk for sale to its customers.

In reference to drug codes 7360 (marihuana) and 7370 (THC), the company plans to bulk manufacture these drugs as synthetics. No other activities for these drug codes are authorized for this registration.

Dated: October 11, 2016.

**Louis J. Milione,**  
Assistant Administrator.

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**DEPARTMENT OF LABOR**

**Wage and Hour Division**

**Agency Information Collection Activities; Comment Request; Information Collections: Report of Construction Contractor’s Wage Rates**

**AGENCY:** Wage and Hour Division, Department of Labor.

**ACTION:** Notice.

**SUMMARY:** The Department of Labor (DOL) is soliciting comments concerning a proposed extension of the information collection request (ICR) titled, “Report of Construction Contractor’s Wage Rates.” This comment request is part of continuing Departmental efforts to reduce paperwork and respondent burden in accordance with the Paperwork Reduction Act of 1995 (PRA), 44 U.S.C. 3501 *et seq.*

This program helps to ensure that requested data can be provided in the desired format, reporting burden (time and financial resources) is minimized, collection instruments are clearly understood, and the impact of collection requirements on respondents can be properly assessed. A copy of the proposed information request can be obtained by contacting the office listed below in the **FOR FURTHER INFORMATION CONTACT** section of this Notice.

**DATES:** Written comments must be submitted to the office listed in the **ADDRESSES** section below on or before December 19, 2016.

**ADDRESSES:** You may submit comments identified by Control Number 1235-0015, by either one of the following methods: *Email: WHDPRAComments@dol.gov; Mail, Hand Delivery, Courier: Division of Regulations, Legislation, and Interpretation, Wage and Hour, U.S. Department of Labor, Room S-3502, 200 Constitution Avenue NW., Washington, DC 20210. Instructions: Please submit one copy of your comments by only one*