

issuance of the proposed registration on or before March 12, 2018. Such persons may also file a written request for a hearing on the application March 12, 2018.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** The Attorney General has delegated his 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 Diversion Control Division ("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 December 29, 2017, Meridian Medical Technologies Inc., 2555 Hermelin Drive, Saint Louis, Missouri 63144 applied to be registered as an importer of the Schedule II control substance for Morphine (9300) the basic class of controlled substance.

The company plans to import the listed controlled substance in finished dosage form for internal analytical purposes only. No other activity for this drug code is authorized for this registration. Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of FDA approved or non-approved finished dosage forms for commercial sale.

Dated: January 31, 2018.

**Susan A. Gibson,**

*Deputy Assistant Administrator.*

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**BILLING CODE 4410-09-P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-392]

#### Importer of Controlled Substances Application: Johnson Matthey Inc.

**ACTION:** Notice of application.

**DATES:** Registered importers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before March 12, 2018. Such persons may also file a written request for a hearing on the application on or before March 12, 2018.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All request for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152.

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In accordance with 21 CFR 1301.34(a), this is notice that on November 9, 2017, Johnson Matthey Inc., Pharmaceutical Materials, 2003

Nolte Drive, West Deptford, NJ 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 coca leaves (9040), raw opium (9600), and poppy straw concentrate (9670) in order to bulk manufacture active pharmaceutical ingredients (API) for distribution to its customers. The company plans to also import thebaine (9333), noroxymorphone (9668), and fentanyl (9801) to use as analytical reference standards, both internally and to be sold to their customers to support testing of Johnson Matthey Inc.'s API's only.

Dated: January 31, 2018.

**Susan A. Gibson,**

*Deputy Assistant Administrator.*

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-392]

#### Importer of Controlled Substances Application: Mylan Technologies Inc.

**ACTION:** Notice of application.

**DATES:** Registered bulk importers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before March 12, 2018. Such persons may also file a written request for a hearing on the application on or before March 12, 2018.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All request for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and