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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-571]

Bulk Manufacturer of Controlled Substances Application: Johnson Matthey Pharmaceutical Materials 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 on or before **[INSERT 60 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER]**.

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 October 31, 2019, Johnson Matthey Pharmaceutical Materials Inc., 25 Patton Road, Devens, Massachusetts 01434 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled Substance	Drug Code	Schedule
Amphetamine	1100	II
Methylphenidate	1724	II
Nabilone	7379	II
Hydrocodone	9193	II
Levorphanol	9220	II
Alfentanil	9737	II
Remifentanil	9739	II

Sufentanil	9740	II

The company plans to utilize this facility to manufacture small quantities of the listed controlled substances in bulk for distribution to its customers as well as to conduct analytical testing in support of the company's primary manufacturing facility in West Deptford, New Jersey.

Dated: December 17, 2019.

William T. McDermott,

Assistant Administrator.

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