



DEPARTMENT OF JUSTICE
Drug Enforcement Administration
Manufacturer of Controlled Substances
Notice of Application
Johnson Matthey, Inc.

Pursuant to § 1301.33(a), Title 21 of the Code of
Federal Regulations (CFR), this is notice that on
September 5, 2013, Johnson Matthey, Inc., Pharmaceuticals
Materials, 900 River Road, Conshohocken, Pennsylvania
19428, made application by renewal to the Drug Enforcement
Administration (DEA) to be registered as a bulk
manufacturer of the following basic classes of controlled
substances:

Drug	Schedule
Gamma Hydroxybutyric Acid (2010)	I
Amphetamine (1100)	II
Methylphenidate (1724)	II
Codeine (9050)	II
Oxycodone (9143)	II

Drug	Schedule
Diphenoxylate (9170)	II
Hydrocodone (9193)	II
Meperidine (9230)	II
Methadone (9250)	II
Methadone intermediate (9254)	II
Morphine (9300)	II
Thebaine (9333)	II

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

The Thebaine (9333) will be used to manufacture other controlled substances for sale in bulk to its customers.

Any other such applicant, and any person who is presently registered with DEA to manufacture such substances, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR § 1301.33(a).

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODW), 8701 Morrisette Drive,

Springfield, Virginia 22152; and must be filed no later than [INSERT DATE 60 DAYS FROM DATE OF PUBLICATION].

Joseph T. Rannazzisi
Deputy Assistant Administrator
Office of Diversion Control
Drug Enforcement Administration

DATED: November 5, 2013

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