Controlled substance	Drug code	Schedule
Meperidine	9230	II.
Meperidine intermediate-B	9233	II.
Methadone	9250	II.
Dextropropoxyphene, bulk (non-dosage forms)	9273	II.
Morphine	9300	II.
Thebaine	9333	II.
Oxymorphone	9652	II.
Alfentanil	9737	II.
Remifentanil	9739	II.
Sufentanil	9740	II.
Carfentanil	9743	II.
Tapentadol	9780	II.
Fentanyl	9801	II.

The company plans to manufacture bulk controlled substances for use in product development of analytical reference standards, for distribution to its customers.

Dated: December 20, 2016.

Louis J. Milione,

Assistant Administrator.

[FR Doc. 2016–31285 Filed 12–27–16; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Registration

ACTION: Notice of registration.

SUMMARY: Registrants listed below have applied for and been granted registration by the Drug Enforcement Administration (DEA) as importers of various classes of schedule I or II controlled substances.

SUPPLEMENTARY INFORMATION:

The companies listed below applied to be registered as importers of various basic classes of controlled substances. Information on previously published notices is listed in the table below. No comments or objections were submitted and no requests for hearing were submitted for these notices.

Company	FR docket	Published
Rhodes Technologies	81 FR 46956	July 19, 2016.
Bellwyck Clinical Services	81 FR 54603	August 16, 2016.
Cerilliant Corporation	81 FR 57933	August 24, 2016.
Noramco, Inc	81 FR 57932	August 24, 2016.
Cody Laboratories, Inc	81 FR 54602	August 16, 2016.
AMRI Rensselaer, Inc	81 FR 54603	August 16, 2016.
ALMAC Clinical Services Incorp (ACSI)	81 FR 54602	August 16, 2016.
Fresenius Kabi USA, LLC	81 FR 54601	August 16, 2016.
Akorn, Inc	81 FR 57935	August 24, 2016.
Actavis Laboratories FL, Inc	81 FR 54602	August 16, 2016.
Unither Manufacturing LLC	81 FR 61250	September 6, 2016.
Cambrex Charles City	81 FR 63222	September 14, 2016.
United States Pharmacopeial Convention	81 FR 63220	September 14, 2016.
R & D Systems, Inc	81 FR 64509	September 20, 2016.
Catalent CTS, LLC	81 FR 66081	September 26, 2016.

The DEA has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of the listed registrants to import the applicable basic classes of schedule I or II controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated each company's maintenance of effective controls against diversion by inspecting and testing each company's physical security systems, verifying each company's compliance with state and local laws, and reviewing each company's background and history.

Therefore, pursuant to 21 U.S.C. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the DEA has granted a registration as an importer for schedule I or II controlled substances to the above listed persons.

Dated: December 19, 2016.

Louis J. Milione,

Assistant Administrator.

[FR Doc. 2016–31273 Filed 12–27–16; 8:45 am]

BILLING CODE 4410–09–P

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 February 27, 2017.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrissette 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