its customers. Dihydromorphine is an intermediate in the manufacture of Hydromorphone and is not for commercial distribution.

Dated: March 14, 2016.

Louis J. Milione,

Deputy Assistant Administrator.

[FR Doc. 2016–06532 Filed 3–22–16; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Manufacturer of Controlled Substances Registration: Cambrex Charles City

ACTION: Notice of registration.

SUMMARY: Cambrex Charles City applied to be registered as a manufacturer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants Cambrex Charles City registration as a manufacturer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated October 13, 2015, and published in the **Federal Register** on October 21, 2015, 80 FR 63835, Cambrex Charles City, 1205 11th Street, Charles City, Iowa 50616 applied to be registered as a manufacturer of certain basic classes of controlled substances. No other comments or objections were submitted for this notice.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Cambrex Charles City to manufacture the basic classes of 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 the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing the company's background and history.

Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR 1301.33, the above-named company is granted registration as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Schedule
Amphetamine (1100)	II
Lisdexamfetamine (1205)	II
Methylphenidate (1724)	II
4-Anilino-N-phenethyl-4-piperidine (ANPP) (8333).	II
Phenylacetone (8501)	II
Cocaine (9041)	II
Codeine (9050)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Hydrocodone (9193)	II
Morphine (9300)	II
Oripavine (9330)	II
Thebaine (9333)	II
Opium extracts (9610)	II
Opium fluid extract (9620)	II
Opium tincture (9630)	II
Opium, powdered (9639)	II
Oxymorphone (9652)	II
Noroxymorphone (9668)	II
Fentanyl (9801)	II

The company plans to manufacture the listed controlled substances in bulk for sale to its customers, for dosage form development, for clinical trials, and for use in stability qualification studies.

Dated: March 14, 2016.

Louis J. Milione,

Deputy Assistant Administrator.

[FR Doc. 2016–06536 Filed 3–22–16; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Registration: Cerilliant Corporation

ACTION: Notice of registration.

SUMMARY: Cerilliant Corporation applied to be registered as an importer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants Cerilliant Corporation registration as an importer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated October 13, 2015, and published in the **Federal Register** on October 21, 2015, 80 FR 63836, Cerilliant Corporation, 811 Paloma Drive, Suite A, Round Rock, Texas 78665–2402 applied to be registered as an importer of certain basic classes of controlled substances. Comments and requests for hearing on applications to import narcotic raw material are not appropriate. 72 FR 3417, (January 25, 2007). No comments or objections were submitted for this notice.

The DEA has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of Cerilliant Corporation to import the basic classes of 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 the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing the 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 above-named company is granted registration as an importer of the following basic classes of controlled substances:

Controlled substance	Schedule
3-Fluoro-N-methylcathinone (3–FMC) (1233)	I
Cathinone (1235)	I
Methcathinone (1237)	I
4-Fluoro-N-methylcathinone (4–FMC) (1238)	I
Pentedrone (*a*-methylaminovalerophenone) (1246)	I
Mephedrone (4-Methyl-N-methylcathinone) (1248)	I
4-Methyl-N-ethylcathinone (4–MEC) (1249)	I
Naphyrone (1258)	I
N-Ethylamphetamine (1475)	I
N,N-Dimethylamphetamine (1480)	I
Fenethylline (1503)	I
Methaqualone (2565)	I
JWH–250 (1-Pentyl-3-(2-methoxyphenylacetyl)indole) (6250)	I
SR–18 (Also known as RCS–8) (1-Cyclohexylethyl-3-(2-methoxyphenylacetyl)indole) (7008)	I