DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-668]

Bulk Manufacturer of Controlled Substances Application: Cambrex Charles City

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic class(es), 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 May 6, 2020, Cambrex Charles City, 1205 11th Street, Charles City, Iowa 50616-3466, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substances:

Controlled Substance	Drug Code	Schedule
Gamma Hydroxybutyric Acid	2010	I
Amphetamine	1100	II
Lisdexamfetamine	1205	II
Methylphenidate	1724	II
ANPP (4-Anilino-N-phenethyl-4-piperidine)	8333	II
Phenylacetone	8501	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 above-listed controlled substances in bulk for conversion to other controlled substances and sales to its customers for dosage form development, clinical trials and use in stability qualification studies. No other activities for these drug codes are authorized for this registration.

William T. McDermott,
Assistant Administrator.

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