projects and programs in the following technology areas: Multi-function Materials, Cyber, Big Data Analytics/ Artificial Intelligence/Machine Learning, Directed Energy Science & Engineering, Advanced Computing and Software Engineering, Autonomous and Unmanned Systems, Sensor Systems, Gun and Projectile Systems, Digital Engineering, Human Systems Integration, Quantum Technologies, Threat Engineering, Mission Engineering and Analysis, Integrated Warfare Systems, Virtualization, Asymmetric Warfare, Manufacturing, Lethality, Surface Offensive & Defensive Engagements, and Launcher Technology.

Suzanne Morris,

Chief, Premerger and Division Statistics Unit, Antitrust Division.

[FR Doc. 2019–24483 Filed 11–8–19; 8:45 am]

BILLING CODE 4410–11–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–541]

Bulk Manufacturer of Controlled Substances Application: Rhodes Technologies

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 January 13, 2020.

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 August 16, 2019, Rhodes Technologies, 498 Washington Street, Coventry, Rhode Island 02816 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Marihuana	7360	I
Tetrahydrocannabinols	7370	I
Dihydromorphine	9145	I
Methylphenidate	1724	II
Codeine	9050	II
Dihydrocodeine	9120	II
Oxycodone	9143	II
Hydromorphone	9150	II
Hydrocodone	9193	II
Levorphanol	9220	II
Morphine	9300	II
Oripavine	9330	II
Thebaine	9333	II
Oxymorphone	9652	II
Noroxymorphone	9668	II
Tapentadol	9780	II
Fentanyl	9801	II

The company plans to manufacture the listed controlled substances in bulk for conversion and sale to finished dosage form manufacturers. In reference to drug codes 7360 (marihuana), and 7370 (tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetic cannabidiol (CBD) and tetrahydrocannabinol (THC). No other activities for drug codes 7360 and 7370 are authorized for this registration.

Dated: October 30, 2019.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2019–24545 Filed 11–8–19; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–539]

Bulk Manufacturer of Controlled Substances Application: Cambrex Charles City

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 January 13, 2020.

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 15, 2019, Cambrex Charles City, 1205 11th Street, Charles City, Iowa 50616–3466 applied to be registered as a bulk manufacturer of the following basic classes 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
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 conversion to other controlled substances and sale to its customers, for dosage form development, for clinical trials, and for use in stability qualification studies.

Dated: October 30, 2019.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2019–24543 Filed 11–8–19; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–534]

Bulk Manufacturer of Controlled Substances Application: Cambridge Isotope Lab

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 January 13, 2020.

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 July 24, 2019 Cambridge Isotope Lab, 50 Frontage Road, Andover, Massachusetts 01810–5413 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances: