DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–579]

Bulk Manufacturer of Controlled Substances Application: Sigma Aldrich Research

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 May 11, 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 December 13, 2019, Sigma Aldrich Research, Biochemicals, Inc., 400–600 Summit Drive, Burlington, Massachusetts 01803 applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substances:

Controlled substance	Drug code	Schedule
Cathinone	1235	I
Mephedrone (4-Methyl-N-methylcathinone)	1248	I
Lysergic acid diethylamide	7315	I
Tetrahydrocannabinols	7370	I
3,4-Methylenedioxymethamphetamine	7405	I
Alpha-methyltryptamine	7432	I
Dimethyltryptamine	7435	I
5-Methoxy-N,N-diisopropyltryptamine	7439	I
N-Benzylpiperazine	7493	I
2-(2,5-Dimethoxyphenyl) ethanamine (2C-H)	7517	I
MDPV (3,4-Methylenedioxypyrovalerone)	7535	I
Methylone (3,4-Methylenedioxy-N-methylcathinone)	7540	I
Heroin	9200	I
Normorphine	9313	I
Norlevorphanol	9634	I
Amphetamine	1100	II
Nabilone	7379	II
Phencyclidine	7471	II
Cocaine	9041	II
Codeine	9050	II
Ecgonine	9180	II
Levorphanol	9220	II
Meperidine	9230	II
Methadone	9250	II
Morphine	9300	II
Thebaine	9333	II
Levo-alphacetylmethadol	9648	II
Noroxymorphone	9668	II
Remifentanil	9739	II
Sufentanil	9740	II
Carfentanil	9743	II
Fentanyl	9801	II

The company plans to manufacture small quantities of the listed controlled substances to make reference standards for distribution to its customers.

Dated: February 10, 2020.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–04831 Filed 3–9–20; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–583]

Bulk Manufacturer of Controlled Substances Application: Siemens Healthcare Diagnostics 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 on or before May 11, 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 December 10, 2019, Siemens Healthcare Diagnostics Inc., 100 GBC Drive, Mailstop 514, Newark, Delaware 19702–2461 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Ecgonine	9180	II

The company plans to produce the listed controlled substance in bulk to be used in the manufacture of DEA exempt products.

Dated: February 11, 2020.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–04832 Filed 3–9–20; 8:45 am]

BILLING CODE 4410–09–P