

Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his

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 respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of, controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been

re delegated to the Assistant Administrator of the DEA Diversion Control Division ("Assistant Administrator") pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on August 25, 2016, Cambridge Isotope Laboratories, 50 Frontage Road, Andover, Massachusetts 01810 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Mephedrone (4-Methyl-N-methylcathinone)	1248	I
Gamma Hydroxybutyric Acid	2010	I
Methaqualone	2565	I
Lysergic acid diethylamide	7315	I
Marihuana	7360	I
Tetrahydrocannabinols	7370	I
3,4-Methylenedioxyamphetamine	7400	I
3,4-Methylenedioxy-N-ethylamphetamine	7404	I
3,4-Methylenedioxymethamphetamine	7405	I
Methylone (3,4-Methylenedioxy-N-methylcathinone)	7540	I
Butylone	7541	I
Heroin	9200	I
Pentobarbital	2270	II
Secobarbital	2315	II
Phencyclidine	7471	II
Cocaine	9041	II
Dihydrocodeine	9120	II
Ecgonine	9180	II
Meperidine	9230	II
Dextropropoxyphene, bulk (non-dosage forms)	9273	II
Fentanyl	9801	II

The company plans to import the listed controlled substances for analytical research, testing and clinical trials.

Dated: April 18, 2017.

Louis J. Milione,
Assistant Administrator.

[FR Doc. 2017-08346 Filed 4-24-17; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Bulk Manufacturer of Controlled Substances Application: Siegfried USA, LLC

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 June 26, 2017.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his 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 respect to the promulgation and

implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been re delegated to the Assistant Administrator of the DEA Diversion Control Division ("Assistant Administrator") pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.33(a), this is notice that on November 22, 2016, Siegfried USA, LLC, 33 Industrial Park Road, Pennsville, New Jersey 08070 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
Dihydromorphine	9145	I
Hydromorphenol	9301	I
Methylphenidate	1724	II
Amobarbital	2125	II
Pentobarbital	2270	II
Secobarbital	2315	II

Controlled substance	Drug code	Schedule
Codeine	9050	II
Oxycodone	9143	II
Hydromorphone	9150	II
Hydrocodone	9193	II
Methadone	9250	II
Methadone intermediate	9254	II
Dextropropoxyphene, bulk (non-dosage forms)	9273	II
Morphine	9300	II
Oripavine	9330	II
Thebaine	9333	II
Opium tincture	9630	II
Oxymorphone	9652	II

The company plans to manufacture the listed controlled substances in bulk for distribution to its customers.

Dated: April 18, 2017.

Louis J. Milione,

Assistant Administrator.

[FR Doc. 2017-08347 Filed 4-24-17; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Bulk Manufacturer of Controlled Substances Application: Sigma Aldrich Research Biochemicals, 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 June 26, 2017.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his 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

respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division ("Assistant Administrator") pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.33(a), this is notice that on December 19, 2016, Sigma Aldrich Research Biochemicals, Inc., 1-3 Strathmore Road, Natick, Massachusetts 01760-2447 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Cathinone	1235	I
Methcathinone	1237	I
Mephedrone (4-Methyl-N-methylcathinone)	1248	I
Aminorex	1585	I
Alpha-ethyltryptamine	7249	I
Lysergic acid diethylamide	7315	I
Tetrahydrocannabinols	7370	I
4-Bromo-2,5-dimethoxyamphetamine	7391	I
4-Bromo-2,5-dimethoxyphenethylamine	7392	I
4-Methyl-2,5-dimethoxyamphetamine	7395	I
2,5-Dimethoxyamphetamine	7396	I
3,4-Methylenedioxyamphetamine	7400	I
N-Hydroxy-3,4-methylenedioxyamphetamine	7402	I
3,4-Methylenedioxy-N-ethylamphetamine	7404	I
3,4-Methylenedioxymethamphetamine	7405	I
Dimethyltryptamine	7435	I
Psilocybin	7437	I
5-Methoxy-N,N-diisopropyltryptamine	7439	I
1-[1-(2-Thienyl)cyclohexyl]piperidine	7470	I
N-Benzylpiperazine	7493	I
MDPV (3,4-Methylenedioxypropylvalerone)	7535	I
Mephedrone (3,4-Methylenedioxy-N-methylcathinone)	7540	I
Heroin	9200	I
Normorphine	9313	I
Norlevorphanol	9634	I
Amphetamine	1100	II
Methamphetamine	1105	II
Nabilone	7379	II
1-Phenylcyclohexylamine	7460	II
Phencyclidine	7471	II
Cocaine	9041	II
Codeine	9050	II
Ecgonine	9180	II