

Dated: July 30, 2012.

Joseph T. Rannazzisi, Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 2012-19191 Filed 8-6-12; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application; Lipomed

Pursuant to Title 21 Code of Federal Regulations 1301.34 (a), this is notice that on June 13, 2012, Lipomed, One Broadway, Cambridge, Massachusetts 02142, made application by letter to the Drug Enforcement Administration (DEA) for registration as an importer of the following basic classes of controlled substances:

Table with 2 columns: Drug, Schedule. Rows include Bufotenine (7433), Diethyltryptamine (7434), 1-Piperidinocyclohexanecarbonitrile (8603).

The company plans to import analytical reference standards for

distribution to its customers for research and analytical purposes.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances listed in schedules I or II, which fall under the authority of section 1002(a)(2)(B) of the Act 21 U.S.C. 952(a)(2)(B) may, in the circumstances set forth in 21 U.S.C. 958(i), file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43, and in such form as prescribed by 21 CFR 1316.47.

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than September 6, 2012

This procedure is to be conducted simultaneously with, and independent of, the procedures described in 21 CFR 1301.34(b), (c), (d), (e), and (f). As noted in a previous notice published in the Federal Register on September 23, 1975, 40 FR 43745-46, all applicants for registration to import a basic class of any controlled substance in schedules I or II are, and will continue to be,

required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, that the requirements for such registration pursuant to 21 U.S.C. 958(a); 21 U.S.C. 823(a); and 21 CFR 1301.34(b), (c), (d), (e), and (f) are satisfied.

Dated: July 30, 2012.

Joseph T. Rannazzisi, Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 2012-19196 Filed 8-6-12; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application; Cerilliant Corporation

Pursuant to Title 21 Code of Federal Regulations 1301.34 (a), this is notice that on July 6, 2012, Cerilliant Corporation, 811 Paloma Drive, Suite A, Round Rock, Texas 78665-2402, made application by renewal to the Drug Enforcement Administration (DEA) for registration as an importer of the following basic classes of controlled substances:

Table with 2 columns: Drug, Schedule. Lists various controlled substances such as Cathinone, Methcathinone, 4-Methyl-N-methylcathinone, N-Ethylamphetamine, N,N-Dimethylamphetamine, Fenethylamine, Gamma Hydroxybutyric Acid, JWH-018, JWH-073, JWH-200, Alpha-ethyltryptamine, Ibogaine, CP-47497, CP-47497 C8 Homologue, Lysergic acid diethylamide, 2,5-Dimethoxy-4-(n)-propylthiophenethylamine, Marihuana, Tetrahydrocannabinols, Mescaline, 3,4,5-Trimethoxyamphetamine, 4-Bromo-2,5-dimethoxyamphetamine, 4-Bromo-2,5-dimethoxyphenethylamine, 4-Methyl-2,5-dimethoxyamphetamine, 2,5-Dimethoxyamphetamine, 3,4-Methylenedioxyamphetamine, 3,4-Methylenedioxy-N-ethylamphetamine, 3,4-Methylenedioxy-methamphetamine, 4-Methoxyamphetamine, 5-Methoxy-N,N-dimethyltryptamine, Alpha-methyltryptamine, Diethyltryptamine, Dimethyltryptamine, Psilocybin, and Psilocyn.

Drug	Schedule
5-Methoxy-N,N-diisopropyltryptamine (7439)	I
N-Benzylpiperazine (7493)	I
3,4-Methylenedioxypropylvalerone (7535)	I
3,4-Methylenedioxy-N-methylcathinone (7540)	I
Desomorphine (9055)	I
Etorphine (except HCl)(9056)	I
Heroin (9200)	I
Morphine-N-oxide (9307)	I
Normorphine (9313)	I
Pholcodine (9314)	I
Dextromoramide (9613)	I
Dipipanone (9622)	I
Racemoramide (9645)	I
Trimeperidine (9646)	I
1-Methyl-4-phenyl-4-propionoxypiperidine (9661)	I
Tilidine (9750)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Methylphenidate (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Phencyclidine (7471)	II
Phenylacetone (8501)	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Benzoyllecgonine (9180)	II
Ethylmorphine (9190)	II
Meperidine (9230)	II
Methadone (9250)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273)	II
Morphine (9300)	II
Oripavine (9330)	II
Thebaine (9333)	II
Levo-alphaacetylmethadol (9648)	II
Oxymorphone (9652)	II
Poppy Straw Concentrate (9670)	II
Alfentanil (9737)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

The company plans to import small quantities of the listed controlled substances for the manufacture of analytical reference standards.

In reference to drug codes 7360 and 7370, the company plans to import a synthetic cannabidiol and a synthetic Tetrahydrocannabinol. No other activity for this drug code is authorized for this registration.

Comments and requests for hearing on applications to import narcotic raw material are not appropriate. 72 FR 3417(2007).

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances listed in schedules I or II, which fall under the authority of section 1002(a)(2)(B) of the Act 21 U.S.C. 952(a)(2)(B) may, in the circumstances set forth in 21 U.S.C. 958(i), file comments or objections to the issuance of the proposed registration and may, at the

same time, file a written request for a hearing on such application pursuant to 21 CFR § 1301.43 and in such form as prescribed by 21 CFR § 1316.47.

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than September 6, 2012.

This procedure is to be conducted simultaneously with, and independent of, the procedures described in 21 CFR § 1301.34(b), (c), (d), (e), and (f). As noted in a previous notice published in the **Federal Register** on September 23, 1975, 40 FR 43745-46, all applicants for registration to import a basic class of any controlled substances in schedules I or II are, and will continue to be, required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement

Administration, that the requirements for such registration pursuant to 21 U.S.C. 958(a); 21 U.S.C. 823(a); and 21 CFR § 1301.34(b), (c), (d), (e), and (f) are satisfied.

Dated: July 30, 2012.

Joseph T. Rannazzisi,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application; Clinical Supplies Management Inc.

Pursuant to Title 21 Code of Federal Regulations 1301.34(a), this is notice that on July 3, 2012, Clinical Supplies