

Dated: January 18, 2011.

**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**

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 25, 2010, AMRI Rensselaer, Inc., 33 Riverside Avenue, Rensselaer, New York 12144, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
Marihuana (7360) .....	I
Tetrahydrocannabinols (7370) .....	I
Amphetamine (1100) .....	II
Lisdexamfetamine (1205) .....	II
Methylphenidate (1724) .....	II
Pentobarbital (2270) .....	II
Meperidine (9230) .....	II
Fentanyl (9801) .....	II

The company plans to manufacture bulk controlled substances for use in product development and for distribution to its customers.

In reference to drug code 7360 (Marihuana), the company plans to bulk manufacture cannabidiol as a synthetic intermediate. This controlled substance will be further synthesized to bulk manufacture a synthetic THC (7370). No other activity for this drug code is authorized for this registration.

Any other such applicant, and any person who is presently registered with DEA to manufacture such substances, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

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 April 4, 2011.

Dated: January 18, 2011.

**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**

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on November 24, 2010, Sigma Aldrich Research Biochemicals, Inc., 1-3 Strathmore Road, Natick, Massachusetts 01760-2447, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following classes of controlled substances:

Drug	Schedule
Cathinone (1235) .....	I
Methcathinone (1237) .....	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
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-Methylenedioxy-N-methylamphetamine (MDMA) (7405) .....	I
Psilocybin (7437) .....	I
5-Methoxy-N,N-diisopropyltryptamine (7439) .....	I
1-[1-(2-Thienyl)cyclohexyl]piperidine (TCP) (7470) .....	I
N-Benzylpiperazine (BZP) (7493) .....	I
Heroin (9200) .....	I
Normorphine (9313) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Nabilone (7379) .....	II
1-Phenylcyclohexylamine (7460) .....	II
Phencyclidine (7471) .....	II
Cocaine (9041) .....	II
Codeine (9050) .....	II
Diprenorphine (9058) .....	II
Ecgonine (9180) .....	II
Levomethorphan (9210) .....	II
Levorphanol (9220) .....	II

Drug	Schedule
Meperidine (9230) .....	II
Metazocine (9240) .....	II
Methadone (9250) .....	II
Morphine (9300) .....	II
Thebaine (9333) .....	II
Levo-alphaacetylmethadol (9648) .....	II
Remifentanyl (9739) .....	II
Carfentanyl (9743) .....	II
Fentanyl (9801) .....	II

The company plans to manufacture reference standards.

Any other such applicant, and any person who is presently registered with DEA to manufacture such substances, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

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 April 4, 2011.

Dated: January 18, 2011.

**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 Registration**

By Notice dated August 3, 2010, and published in the **Federal Register** on September 1, 2010, 75 FR 53719, Cambridge Isotope Lab, 50 Frontage Road, Andover, Massachusetts 01810, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Morphine (9300), a basic class of controlled substance listed in schedule II.

The company plans to utilize small quantities of the listed controlled substance in the preparation of analytical standards.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Cambridge Isotope Lab to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has