

By order of the Commission.

Donna R. Koehnke,
Secretary.

[FR Doc. 97-33210 Filed 12-18-97; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated June 23, 1997, and published in the **Federal Register** on July 10, 1997 (62 FR 37077), Applied Science Labs, Division of Alltech Associates, Inc., 2701 Carolean Industrial Drive, P.O. Box 440, State College, Pennsylvania 16801, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Methcathinone (1237)	I
N-Ethylamphetamine (1475)	I
N,N-Dimethylamphetamine (1480)	I
4-Methylaminorex (cis isomer) (1590)	I
Lysergic acid diethylamide (7315)	I
Mescaline (7381)	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
N-Ethyl-1-phenylcyclohexylamine (7455)	I
1-(1-Phenylcyclohexyl) pyrrolidine (7458)	I
1-[1-(2-Thienyl) cyclohexyl]piperidine (7470)	I
Dihydromorphine (9145)	II
1-Phenylcyclohexylamine (7460)	II
Phencyclidine (7471)	II
Phenylacetone (8501)	II
1-Piperidinocyclohexanecarbonitrile (8603)	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Benzoylcegonine (9180)	II
Morphine (9300)	II
Oxymorphone (9652)	II
Noroxymorphone (9668)	II

The firm plans to manufacture small quantities of the listed controlled substances for reference standards.

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Applied Science Labs to manufacture the listed controlled substances is consistent with the public

interest at this time. Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: November 28, 1997.

John H. King,

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

[FR Doc. 97-33113 Filed 12-18-97; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on August 25, 1997, Cambridge Isotope Lab, 50 Frontage Road, Andover, Massachusetts 01810, made application by renewal, which was received for processing on November 4, 1997, to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Methaqualone (2565)	I
Dimethyltryptamine (7435)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Phencyclidine (7471)	II
Phenylacetone (8501)	II
Cocaine (9041)	II
Codeine (9050)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Benzoylcegonine (9180)	II
Methadone (9250)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273)	II
Morphine (9300)	II
Fentanyl (9801)	II

The firm plans to manufacture small quantities of the above listed controlled substances for isotope labeled standards for drug analysis.

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.

Any such comments or objections may be addressed, in quintuplicate, to

the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than February 17, 1998.

Dated: November 28, 1997.

John H. King,

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

[FR Doc. 97-33115 Filed 12-18-97; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated June 9, 1997, and published in **Federal Register** on June 17, 1997, (62 FR 32824), Damocles10, 3529 Lincoln Highway, Thorndale, Pennsylvania 19372, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Heroin (9200)	I
Codeine (9050)	II
Hydromorphone (9150)	II
Hydrocodone (9193)	II
Morphine (9300)	II

The firm plans to manufacture the listed controlled substances for the purpose of deuterium labeled internal standards for distribution to analytical laboratories.

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Damocles10 to manufacture the listed controlled substances is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. 823 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: December 1, 1997.

John H. King,

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

[FR Doc. 97-33116 Filed 12-18-97; 8:45 am]

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