

American Radiolabeled Chemical, Inc., on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, verification of the company's compliance with state and local laws, and a review of the company's background and history. 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 class of controlled substance listed above is granted.

Dated: August 18, 2000.

**John H. King,**

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

[FR Doc. 00-22691 Filed 9-5-00; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled Substances; Notice of Application**

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on June 13, 2000, 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) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Methacathinone (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

Drug	Schedule
1-[1-(2-Thienyl)cyclohexyl]piperidine (7470).	I
Dihydromorphine (9145) .....	I
Normorphine (9313) .....	I
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
Noroxymorphone (9668) .....	II

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

Any other such applicant and any person who is presently registered with DEA to manufacture such substance 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, DC 20537, Attention: DEA Federal Register Representative (CCGR), and must be filed no later than November 6, 2000.

Dated: August 21, 2000.

**John H. King,**

*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 April 18, 2000, and published in the **Federal Register** on April 28, 2000, (65 FR 24985), Eli-Elsohly Laboratories, Inc., 5 Industrial Park Drive, Oxford, Mississippi 38655, 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
Tetrahydrocannabinols (7370) .....	I
Dihydromorphine (9145) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Cocaine (9041) .....	II
Codeine (9050) .....	II

Drug	Schedule
Dihydrocodeine (9120) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Benzoylcegonine (9180) .....	II
Hydrocodone (9193) .....	II
Morphine (9300) .....	II

The firm plans to bulk manufacture non-deuterated controlled substances for use as analytical standards and deuterated controlled substances for use as internal standards.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Eli-Elsohly Laboratories, Inc., to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Eli-Elsohly Laboratories, Inc., on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, verification of the company's compliance with state and local laws, and a review of the company's background and history. 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: August 21, 2000.

**John H. King,**

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

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Importation of Controlled Substances; Notice of Application**

Pursuant to Section 1008 of the Controlled Substances Import and Export Act (21 U.S.C. 958(i)), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in Schedule I or II and prior to issuing a regulation under Section 1002(a) authorizing the importation of such a substance, provide manufacturers holding registrations for