

2000, Noramco, Inc., 1400 Olympic Drive, Athens, Georgia 30601, 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
Amphetamine (1100)	II
Codeine (9050)	II
Oxycodone (9143)	II
Hydrocodone (9193)	II
Morphine (9300)	II
Thebaine (9333)	II
Fentanyl (9801)	II

The firm plans to support its other manufacturing facility with manufacturing and analytical testing.

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 November 24, 2000.

Dated: September 6, 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 the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with Section 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on August 3, 2000, Roche Diagnostics Corporation, 9115 Hague

Road, Indianapolis, Indiana 46250, made application by letter to the Drug Enforcement Administration to be registered as an importer of alpramethadol (9605) a basic class of controlled substance listed in Schedule I.

The firm plans to import alpramethadol to manufacture diagnostic products for distribution to its customers.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substance may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.43 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections or requests for a hearing 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 (CCR), and must be filed no later than October 25, 2000.

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 at 40 FR 43745-46 (September 23, 1975), all applicants for registration to import a basic class of any controlled substance in Schedule 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(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: September 6, 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 May 12, 2000, and published in the **Federal Register** on June 2, 2000, (65 FR 35397), Sigma Aldrich Research Biochemicals, Inc.,

Attn: Richard Milius, 1-3 Strathmore Road, Natick, Massachusetts 01760, 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
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-dimethoxyam- phetamine (7391)	I
4-Bromo-2, 5-dimethoxyphen- ethylamine (7392)	I
2, 5-Dimethoxyamphetamine (7396).	I
3, 4-Methylenedioxyamphetamine (7400).	I
N-Hydroxy-3, 4-methylenedioxy- amphetamine (7402)	I
3, 4-Methylenedioxy-N-ethylam- phetamine (7404)	I
3,4- Methylenedioxymethamphetam- ine (7405).	I
Psilocybin (7437)	I
1-[1- (2-Thienyl)cyclo- hexyl]piperidine (7470)	I
Heroin (9200)	I
Normorphine (9313)	II
Amphetamine (1100)	II
Methamphetamine (1105)	II
Pentobarbital (2270)	II
Phenylcyclohexylamine (7460)	II
Phencyclidine (7471)	II
Cocaine (9041)	II
Codenine (9050)	II
Diprenorphine (9058)	II
Benzoylcgonine (9180)	II
Levomethorphan (9210)	II
Levorphanol (9220)	II
Meperidine (9230)	II
Metazocine (924)	II
Methadone (9250)	II
Morphine (9300)	II
Thebaine (9333)	II
Levo-alphaacetylmethadol (LAAM) (9648).	II
Fentanyl (9801)	II

The firm plans to manufacture the listed controlled substances for laboratory reference standards and neurochemicals.

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Sigma Aldrich Research Biochemicals, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Sigma Aldrich Research Biochemicals, Inc. on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included