

moving all bulk manufacturing of methylphenidate to the above location.

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 (60 days from publication).

Dated: July 29, 1997.

John H. King,

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

[FR Doc. 97-22564 Filed 8-25-97; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated April 24, 1997, and published in the **Federal Register** on May 21, 1997 (62 FR 27770), Research Biochemicals, Inc., Limited Partnership, Attn: Richard Milius, 1-3 Strathmore Road, Natick, Massachusetts 01760, made application to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Methaqualone (2565)	I
Ibogaine (7260)	I
Tetrahydrocannabinols (7370)	I
Bufotenine (7433)	I
Dimethyltryptamine (7435)	I
Etorphine (except Hcl) (9056)	I
Methylphenidate (1724)	II
Pentobarbital (2270)	II
Diprenorphine (9058)	II
Etorphine Hydrochloride (9059) ...	II
Diphenoxylate (9170)	II
Metazocine (9240)	II
Methadone (9250)	II
Fentanyl (9801)	II

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 Research Biochemicals to

import the listed controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971, at this time. Therefore, pursuant to Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, Section 1301.34, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: July 21, 1997.

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 24, 1997, and published in the **Federal Register** on May 21, 1997, (62 FR 27776), Research Biochemicals, Limited Partnership, One Strathmore Road, Natick, Massachusetts 01760, made application 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
Alpha-Ethyltryptamine (7249)	I
Lysergic acid diethylamide (7315)	I
2,5-Dimethoxyamphetamine (7396).	I
3,4-Methylenedioxymethamphetamine (7405). Dimethyltryptamine (7435).	I
1-[1-(2-Thienyl)cyclohexyl]piperidine (7470).	I
Heroin (9200)	I
Normorphine (9313)	I
Phencyclidine (7471)	II
Benzoylcegonine (9180)	II

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Research Biochemicals 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 C.F.R. §§ 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: July 29, 1997.

John H. King,

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

[FR Doc. 97-22566 Filed 8-25-97; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated April 15, 1997, and published in the **Federal Register** on May 12, 1997, (62 FR 25972), Research Triangle Institute, Kenneth H. Davis, Jr., Hermann Building, East Institute Drive, P.O. Box 12194, Research Triangle Park, North Carolina 27709, 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
Marihuana (7360)	I
Cocaine (9041)	II

DEA has considered the factors in Title 21, United States Code, Section 823(a), as well as information provided by other bulk manufacturers, and determined that the registration of Research Triangle Institute 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 C.F.R. §§ 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: July 29, 1997.

John H. King,

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

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