controlled substances listed above is granted.

Dated: March 11, 2003.

Laura M. Nagel,

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

[FR Doc. 03–7834 Filed 4–1–03; 8:45 am]

BILLING CODE 4410–09–M

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 November 18, 2002, Novartis Pharmaceuticals Corporation, Attn: Security Department, Building 103, Room 335, 59 Route 10, East Hanover, New Jersey 07936, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic class of controlled substance listed below:

Drug	Schedule
Methylphenidate (1724)	II

The firm plans to produce bulk product and finished dosage units for distribution to its customers.

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: Drug Operations Section, Domestic Drug Unit (ODOD) and must be filed no later than June 2, 2003.

Dated: March 11, 2003.

Laura M. Nagel,

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

[FR Doc. 03–7829 Filed 4–1–03; 8:45 am]

BILLING CODE 4410–09–M

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 April 11, 2002, Organichem Corporation, 33 Riverside Avenue, Rensselaer, New York, 12144, made application by renewal to the Drug Enforcement Administration for registration as a bulk manufacturer of the basic classes of controlled substance listed below:

Drug	Schedule
Amphetamine (1100)	II
Pentobarbital (2270)	II
Methylphenidate (1724)	II
Meperidine (9230)	II

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: Drug Operations Section, Domestic Drug Unit (ODOD), and must be filed no later than June 2, 2003.

Dated: March 11, 2003

Laura M. Nagel,

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

[FR Doc. 03–7831 Filed 4–1–03; 8:45 am]

BILLING CODE 4410–09–M

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 July 29, 2002, Penick Corporation, 158 Mount Olivet Avenue, Newark, New Jersey 07114, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Cocaine (9040)	II
Ecgonine (9180)	II

The firm plans to manufacture the listed controlled substance for the manufacture of a non-controlled substance flavor extract.

Any other such applicant and any person who is presently registered with the 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 Administrator, United States Department of Justice, Washington, DC 20537, Attention: Drug Operations Section, Domestic Drug Unit (ODOD) and must be filed no later than June 2, 2003.

Dated: March 11, 2003.

Laura M. Nagel,

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

[FR Doc. 03–7828 Filed 4–1–03; 8:45 am]

BILLING CODE 4410–09–M

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 January 9, 2003, Rhodes Technologies, 498 Washington Street, Coventry, Rhode Island 02816, made application by renewal and by letters dated January 28, 2003 and February 26, 2003, to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic class of Schedule I and II controlled substances listed below:

Drug	Schedule
Codeine (9050)	II
Oxycodone (9143)	II
Thebaine (9333)	II
Methylphenidate (1724)	II
Tetrahydrocannabinols (7370)	I
Hydromorphone (9150)	II
Hydrocodone (9193)	II
Fentanyl (9801)	II
Noroxymorphone (9668)	II
Dihydrocodeine (9120)	II

The firm plans to produce bulk products for conversion and distribution to its customers.

Any other such applicant and any person who is presently registered with