

Dated: December 14, 1998.

**John H. King,**

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

[FR Doc. 98-33886 Filed 12-22-98; 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 May 17, 1998, Eli-Elsohly Laboratories, Inc., Mahmoud A. Elsohly, PhD, 5 Industrial Park Drive, Oxford, Mississippi 38655, 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
Tetrahydrocannabinols (7370) ....	I
Amphetamine (1100) .....	II
Methamphetamine (1150) .....	II
Cocaine (9041) .....	II
Codeine (9050) .....	II
Dihydrocodeine (9120) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Benzoylcegonine (9180) .....	II
Hydrocodone (9193) .....	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.

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, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than February 22, 1999.

Dated: December 14, 1998.

**John H. King,**

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

[FR Doc. 98-33881 Filed 12-27-98; 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 October 12, 1998, High Standard Products, 1100 W. Florence Avenue, #B, Inglewood, California 90301, 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
Methaqualone (2565) .....	I
Lysergic acid diethylamide (7315) .....	I
Tetrahydrocannabinols (7370) ..	I
3,4-Methylenedioxyampheta- mine (7400) .....	I
3,4-Methylenedioxy-N- ethylamphetamine (7404) .....	I
3,4-Methylenedioxy- m- thamphetamine (7405) .....	I
4-Methoxyamphetamine (7411)	I
Heroin (9200) .....	I
3-Methylfentanyl (9813) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Secobarbital (2315) .....	II
Phencyclidine (7471) .....	II
Cocaine (9041) .....	II
Codeine (9050) .....	II
Hydromorphone (9150) .....	II
Diphenoxylate (9170) .....	II
Hydrocodone (9193) .....	II
Methadone (9250) .....	II
Morphine (9300) .....	II
Fentanyl (9801) .....	II

The firm plans to manufacture analytical reference standards.

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 22, 1999.

Dated: December 2, 1998.

**John H. King,**

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

[FR Doc. 98-33889 Filed 12-23-98; 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 August 4, 1998, Irix Pharmaceuticals, Inc., 101 Five Star Way, Florence, South Carolina 29501, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of methylphenidate (1724), a basic class of controlled substance listed in Schedule II.

The firm place to manufacture methylphenidate for demonstration purposes and for dosage form development and stability studies.

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, D.C. 20537, Attention: DEA Federal Register Representatives (CCR), and must be filed no later than February 22, 1999.