

DEA has considered the factors in title 21, United States Code, section 823(a) and determined that the registration of Chirex Technology Center, Inc., DBA Chirex Cauldron to manufacture amphetamine is consistent with the public interest at this time. DEA has investigated the firm to ensure that the company's registration is consistent with the public interest. The investigation 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: June 7, 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 March 30, 2000, Radian International LLC, 14050 Summit Drive #121, P.O. Box 201088, Austin, Texas 78720-1088, made application by renewal to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Cathinone (1235) .....	I

Drug	Schedule
Methcathinone (1237) .....	I
N-Ethylamphetamine (1475) .....	I
gamma hydroxybutyric acid (2010).	I
Ibogaine (7260) .....	I
Tetrahydrocannabinols (7370) .....	I
Mescaline (7381) .....	I
4-Bromo-2,5-dimethoxyamphetamine (7391).	I
4-Bromo-2,5-dimethoxyphenethylamine (7392).	I
4-Methyl-2,5-dimethoxyamphetamine (7395).	I
2,5-Dimethoxyamphetamine (7396).	I
3,4-Methylenedioxyamphetamine (7400).	I
3,4-Methylenedioxy-N-ethylamphetamine (7404).	I
3,4-Methylenedioxymethamphetamine (7405).	I
4-Methoxyamphetamine (7411) ..	I
Psilocybin (7437) .....	I
Psilocyn (7438) .....	I
Ethorphine (except HC1) (9056) ..	I
Heroin (9200) .....	I
Pholcodine (9314) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Amobarbital (2125) .....	II
Pentobarbital (2270) .....	II
Cocaine (9041) .....	II
Codeine (9050) .....	II
Dihydrocodeine (9120) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Benzoyllecgonine (9180) .....	II
Ethylmorphine (9190) .....	II
Meperidine (9230) .....	II
Methadone (9250) .....	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II
Morphine (9300) .....	II
Thebaine (9333) .....	II
levo-alphaacetylmethadol (9648) ..	II
Oxymorphone (9652) .....	II

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

Any manufacturer holding, or applying for, registration as a bulk manufacturer of these basic classes of controlled substances 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, D.C. 20537, Attention: DEA Federal Register

Representative (CCR), and must be filed no later than July 24, 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 basic classes of any controlled substances 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 1311.42(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: June 8, 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 Application

Pursuant to section 1301.33(a) of title 21 of the Code of Federal Regulations (CFR), this is notice that on December 10, 1999, Salisbury Chemicals, Inc., 1205 11th Street, Charles City, Iowa 50616-3466, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of amphetamine (1100) and by letter dated March 14, 2000, for registration to bulk manufacture methylphenidate (1724), basic classes of controlled substances listed in Schedule II.

The firm plans to manufacture amphetamine and methylphenidate for distribution as bulk product.

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 Representative (CCR), and must be filed no later than August 21, 2000.