

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated July 26, 2006, and published in the Federal Register on August 2, 2006, (71 FR 43814-43815), Siegfried (USA) Inc., 33 Industrial Park Road, Pennsville, New Jersey 08070, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Hydromorphone (9150), a basic class of controlled substance listed in schedule II.

The company plans to manufacture in bulk for distribution to its customers.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Siegfried (USA), Inc. to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Siegfried (USA), Inc. to ensure that the company's registration is consistent with the public interest. The investigation has 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 in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic class of controlled substance listed.

Dated: May 10, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7-9628 Filed 5-17-07; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated February 16, 2007, and published in the Federal Register on February 27, 2007 (72 FR 8793), Sigma Aldrich Manufacturing LLC., Subsidiary of Sigma-Aldrich Company, 3500 Dekalb Street, St. Louis, Missouri 63118, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed in schedule I and II:

Table with 2 columns: Drug and Schedule. Lists various controlled substances such as Cathinone, Methcathinone, Aminorex, Gamma Hydroxybutyric Acid, Methaqualone, Ibogaine, Lysergic acid diethylamide, Marihuana, Tetrahydrocannabinols, Mescaline, 4-Bromo-2,5-dimethoxyamphetamine, 4-Bromo-2,5-dimethoxyphenethylamine, 4-Methyl-2,5-dimethoxyamphetamine, 2,5-Dimethoxyamphetamine, 3,4-Methylenedioxyamphetamine, N-Hydroxy-3,4-methylenedioxyamphetamine, 3,4-Methylenedioxy-N-ethylamphetamine, 3,4-Methylenedioxy-N-ethylamphetamine (MDMA), 4-Methoxyamphetamine, Bufotenine, Diethyltryptamine, Dimethyltryptamine, Psilocybin, Psilocyn, N-Ethyl-1-phenylcyclohexylamine, N-Benzylpiperazine (BZP), Trifluoromethylphenyl Piperazine, Heroin, Normorphine, Etonitazene, Amphetamine, Methamphetamine, Methylphenidate, Amobarbital, Pentobarbital, Secobarbital, Glutethimide, Nabilone, Phencyclidine, Cocaine, Codeine, Diprenorphine, Oxycodone, Hydromorphone, Diphenoxylate, Ecgonine, Ethylmorphine, Hydrocodone, Levorphanol, Meperidine, Methadone, Dextropropoxyphene, Morphine, Thebaine, Opium powdered, Oxymorphone, Fentanyl.

The company plans to import the listed controlled substances for sale to research facilities for drug testing and analysis.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and 952(a) and determined that the registration of Sigma Aldrich Manufacturing LLC. to import the basic classes of 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. DEA has investigated Sigma Aldrich Manufacturing LLC. to ensure that the company's registration is consistent with the public interest. The investigation has 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. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above named company is granted registration as an importer of the basic classes of controlled substances listed.

Dated: May 14, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7-9604 Filed 5-17-07; 8:45 am]

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DEPARTMENT OF JUSTICE

Foreign Claims Settlement Commission

[F.C.S.C. Meeting Notice No. 4-07]

Sunshine Act Meeting

The Foreign Claims Settlement Commission, pursuant to its regulations (45 CFR Part 504) and the Government in the Sunshine Act (5 U.S.C. 552b), hereby gives notice in regard to the scheduling of meetings for the transaction of Commission business and other matters specified, as follows:

DATE AND TIME: Thursday, May 31, 2007, at 1 p.m.

SUBJECT MATTER: Issuance of Proposed Decisions and Amended Proposed Decisions in claims against Albania.

STATUS: Open.

All meetings are held at the Foreign claims Settlement Commission, 600 E Street, NW., Washington, DC. Requests for information, or advance notices of intention to observe an open meeting, may be directed to: Administrative