

Drug	Schedule
4-Bromo-2,5-dimethoxyampheta- mine (7391).	I
4-Bromo-2,5-dimethoxyphenethylami- ne (7392).	I
2,5-Dimethoxyampheta- mine (7396).	I
3,4-Methylenedioxyampheta- mine (7400).	I
N-Hydroxy-3,4-methylenedioxyampheta- mine (7402).	I
3,4-Methylenedioxy-N- ethylamphetamine (7404).	I
3,4-Methylenedioxymeth- amphetamine (MDMA) (7405).	I
1-[1-(2-Thienyl)cyclohexyl]piperidine (TCP) (7470).	I
1-Benzylpiperazine (BZP) (7493).	I
Heroin (9200) .....	I
Normorphine (9313) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Nabilone (7379) .....	II
1-Phenylcyclohexylamine (7460).	II
Phencyclidine (7471) .....	II
Cocaine (9041) .....	II
Codeine (9050) .....	II
Diprenorphine (9058) .....	II
Ecgonine (9180) .....	II
Levomethorphan (9210) ....	II
Levorphanol (9220) .....	II
Meperidine (9230) .....	II
Metazocine (9240) .....	II
Methadone (9250) .....	II
Morphine (9300) .....	II
Thebaine (9333) .....	II
Levo-alphaacetylmethadol (9648).	II
Carfentanil (9743) .....	II
Fentanyl (9801) .....	II

The company plans to manufacture reference standards. 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 Sigma Aldrich Research BioChemicals, Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Sigma Aldrich Research BioChemicals, 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 classes of controlled substances listed.

Dated: November 28, 2006.  
**Joseph T. Rannazzisi,**  
*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*  
 [FR Doc. E6-20737 Filed 12-6-06; 8:45 am]  
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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Importer of Controlled Substances; Notice of Registration**

By Notice dated July 10, 2006, and published in the **Federal Register** on July 24, 2006, (71 FR 41839-41840), Stepan Company, Natural Products Department, 100 W. Hunter Avenue, Maywood, New Jersey 07607, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Coca Leaves (9040), a basic class of controlled substance listed in schedule II.

The company plans to import the listed controlled substance for the manufacture of bulk controlled substances and distribution to its customers.

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 Stepan Company to import this basic class of controlled substance 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 Stepan Company 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 system, 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 class of controlled substance listed.

Dated: November 28, 2006.  
**Joseph T. Rannazzisi,**  
*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*  
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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Importer of Controlled Substances; Notice of Registration**

By Notice dated August 15, 2006 and published in the **Federal Register** on August 22, 2006, (71 FR 48948), Wildlife Laboratories, Inc., 1401 Duff Drive, Suite 400, Fort Collins, Colorado 80524, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Etorphine Hydrochloride (9059), a basic class of controlled substance listed in schedule II.

The company plans to import the listed controlled substance for sale to its customers.

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 Wildlife Laboratories, Inc. to import the basic class of controlled substance 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 Wildlife Laboratories, 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. 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 class of controlled substance listed.

Dated: November 28, 2006.  
**Joseph T. Rannazzisi,**  
*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*  
 [FR Doc. E6-20741 Filed 12-6-06; 8:45 am]  
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**DEPARTMENT OF JUSTICE**

**Office of Justice Programs**

[OMB Number 1121-0309]

**Agency Information Collection Activities: Proposed Collection; Comments Requested**

**ACTION:** 60-Day Notice of Information Collection Under Review: International Terrorism Victim Expense Reimbursement Program Application.