

Dated: October 16, 2009.

Joseph T. Rannazzisi,
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 § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on April 9, 2009, Aldrich Chemical Company Inc., DBA Isotec, 3858 Benner Road, Miamisburg, Ohio 45342-4304, made application by renewal to the Drug Enforcement Administration (DEA) as a bulk manufacturer of the basic classes of controlled substances listed in schedules I and II:

Drug	Schedule
Gamma Hydroxybutyric Acid (2010).	I
Methaqualone (2565)	I
Ibogaine (7260)	I
Tetrahydrocannabinols (7370)	I
2,5-Dimethoxyamphetamine (7396).	I
Psilocyn (7438)	I
Normorphine (9313)	I
Acetylmethadol (9601)	I
Alphacetylmethadol except levo-alphaacetylmethadol (9603).	I
Normethadone (9635)	I
Norpipanone (9636)	I
3-Methylfentanyl (9813)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Methylphenidate (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
1-Phenylcyclohexylamine (7460)	II
Phencyclidine (7471)	II
Phenylacetone (8501)	II
1-Piperidinocyclohexane-carbonitrile (8603).	II
Cocaine (9041)	II
Codeine (9050)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Benzoyllecgonine (9180)	II
Ethylmorphine (9190)	II
Hydrocodone (9193)	II
Isomethadone (9226)	II
Meperidine (9230)	II
Meperidine intermediate-A (9232)	II
Meperidine intermediate-B (9233)	II
Methadone (9250)	II
Methadone intermediate (9254) ...	II
Dextropropoxyphene, bulk, (non-dosage forms) (9273).	II
Morphine (9300)	II
Thebaine (9333)	II
Levo-alphaacetylmethadol (9648) ..	II

Drug	Schedule
Oxymorphone (9652)	II

The company plans to manufacture small quantities of the listed controlled substances to produce isotope labeled standards for drug testing and analysis. 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 pursuant to 21 CFR 1301.33(a).

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than December 28, 2009.

Dated: October 20, 2009.

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

[FR Doc. E9-25893 Filed 10-27-09; 8:45 am]

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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 September 10, 2009, Johnson Matthey Inc., Custom Pharmaceuticals Department, 2003 Nolte Drive, West Deptford, New Jersey 08066-1742, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in schedules I and II:

Drug	Schedule
Gamma Hydroxybutyric Acid (2010).	I
Tetrahydrocannabinols (7370)	I
Dihydromorphine (9145)	I
Difenoxin (9168)	I
Propiram (9649)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Lisdexamfetamine (1205)	II
Methylphenidate (1724)	II
Nabilone (7379)	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Ecgonine (9180)	II
Hydrocodone (9193)	II

Drug	Schedule
Meperidine (9230)	II
Methadone (9250)	II
Methadone intermediate (9254) ...	II
Morphine (9300)	II
Thebaine (9333)	II
Oxymorphone (9652)	II
Noroxymorphone (9668)	II
Alfentanil (9737)	II
Remifentanil (9739)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

The company plans to manufacture the listed controlled substances in bulk for sale to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such a controlled substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than December 28, 2009.

Dated: October 21, 2009.

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

[FR Doc. E9-26002 Filed 10-27-09; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

**Manufacturer of Controlled
Substances; Notice of Registration**

By Notice dated June 15, 2009, and published in the **Federal Register** on June 23, 2009, (74 FR 29720), Noramco Inc., Division of Ortho-McNeil, Inc., 1440 Olympic Drive, Athens, Georgia 30601, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Noroxymorphone (9668), a basic class of controlled substance listed in schedule II.

The company plans to bulk manufacture the above listed controlled substance 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 Noramco, Inc. to manufacture the listed basic class of controlled substance is consistent with the public interest at

this time. DEA has investigated Noramco, 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: October 16, 2009.

Joseph T. Rannazzisi,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.
 [FR Doc. E9-25887 Filed 10-27-09; 8:45 am]
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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated July 1, 2009, and published in the **Federal Register** on July 13, 2009, (74 FR 33476), Organix Inc., 240 Salem Street, Woburn, Massachusetts 01801, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Marijuana (7360), a basic class of controlled substance listed in schedule I.

The company plans to manufacture a synthetic cannabinol in bulk for sale to its customers for research purposes. No other activity for this drug code is authorized for this registration.

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 Organix Inc. to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Organix 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: October 20, 2009.

Joseph T. Rannazzisi,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.
 [FR Doc. E9-25892 Filed 10-27-09; 8:45 am]
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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances Notice of Registration

By Notice dated June 22, 2009, and published in the **Federal Register** on June 26, 2009, (74 FR 30621), Wildlife Laboratories Inc., 1401 Duff Drive, Suite 400, Fort Collins, Colorado 80524, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Carfentanil (9743), a basic class of controlled substance listed in schedule II.

The company will manufacture the above listed controlled substance for sale to veterinary pharmacies, zoos, and for other animal and wildlife applications.

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 Wildlife Laboratories Inc. to manufacture the listed basic class(es) of controlled substance(s) is consistent with the public interest 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. 823, and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic class(es) of controlled substance(s) listed.

Dated: October 16, 2009.

Joseph T. Rannazzisi,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.
 [FR Doc. E9-25886 Filed 10-27-09; 8:45 am]
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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated February 5, 2009 and published in the **Federal Register** on February 11, 2009, (74 FR 6921), 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 the basic classes of controlled substances listed in schedules I and II:

Drug	Schedule
Dihydromorphine (9145)	I
Amphetamine (1100)	II
Methylphenidate (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Glutethimide (2550)	II
Codeine (9050)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Hydrocodone (9193)	II
Methadone (9250)	II
Methadone intermediate (9254) ...	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II
Morphine (9300)	II
Oripavine (9330)	II
Oxymorphone (9652)	II

The company plans to manufacture the listed controlled substances 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 classes of controlled substances 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 classes of controlled substances listed.