

States obligations under international treaties, conventions, or protocols in effect on May 1, 1971, 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. 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 substances listed.

Dated: July 19, 2005.

William J. Walker,

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 March 25, 2005 and published in the **Federal Register** on April 4 2005, (70 FR 17126), Roche Diagnostics Operations Inc., Attention: Regulatory Compliance, 9115 Hague Road, Indianapolis, Indiana 46250, 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 Schedules I and II:

Drug	Schedule
Lysergic acid diethylamide (7315)	I
Tetrahydrocannabinols (7370)	I
Cocaine (9041)	II
Ecgonine (9180)	II
Methadone (9250)	II
Morphine (9300)	II
Alphamethadol (9605)	II

The company plans to import the listed controlled substances for the manufacture of diagnostic products 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 952(a) and determined that the registration of Roche Diagnostics Operations Inc. 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 Roche Diagnostics Operations 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 classes of controlled substances listed.

Dated: July 19, 2005.

William J. Walker,

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 Registration

By Notice dated February 23, 2005, and published in the **Federal Register** on March 4, 2005, (70 FR 10683), Siegfried (USA), Inc., 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 class of controlled substances listed in Schedule II:

Drug	Schedule
Amphetamine (1100)	II
Methylphenidate (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Glutethimide (2550)	II
Codeine (9050)	II
Hydrocodone (9193)	II
Methadone (9250)	II
Methadone Intermediate (9254) ...	II
Dextropropoxyphene, bulk (non-dosage form) (9273).	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 class 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 class of controlled substances listed.

Dated: July 19, 2005.

William J. Walker,

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated March 29, 2005 and published in the **Federal Register** on April 6, 2005, (70 FR 17474), 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. Sections 823(a) and 952(a) and determined that the registration of Stepan Company 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 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 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. Sections 952(a) and 958(a), and in