DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated October 12, 2001, and published in the **Federal Register** on October 25, 2001, (66 FR 54033), Noramco Inc., 1440 Olympic Drive, Athens, Georgia 30601, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of phenylacetone (8501), basic class of controlled substance listed in Schedule II.

The firm plans to import phenylacetone for the production of amphetamine.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, section 823(a) and determined that the registration of Noramco Inc., 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 Noramco Inc., 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 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 section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, § 1301.34, the above firm is granted registration as an importer of the basic class of controlled substance listed above.

Dated: March 27, 2002.

Laura M. Nagel,

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

[FR Doc. 02–8664 Filed 4–9–02; 8:45 am]

BILLING CODE 4410–09–M

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 September 6, 2001, Organix, Inc., 240 Salem Street, Woburn, Massachusetts 01801, made application by renewal to the Drug Enforcement Administration (DEA) of registration as a bulk manufacturer of cocaine (9041), a basic class of controlled substance listed in Schedule II.

The firm plans to manufacture a derivative of cocaine in gram quantities for validation of synthetic procedures.

Any other such applicant and any person who is presently registered with DEA to manufacturer such substances 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 Assistance Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than June 10, 2002.

Dated: March 27, 2002.

Laura M. Nagel,

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

[FR Doc. 02–8666 Filed 4–9–02; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated August 9, 2001, and published in the **Federal Register** on August 10, 2001, (66 FR 42239), Pressure Chemical Company, 3419 Smallman Street, Pittsburgh, Pennsylvania 15201, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of 2,5-dimethoxyamphetamine (7396), a basic class of controlled substance listed in Schedule I.

The firm plans to bulk manufacture 2,5-dimethoxyamphetamine for distribution to its customers.

No comment or objections have been received. DEA has considered the factors in Title 21, United States Code, section 823(a) and determined that the registration of Pressure Chemical Company to manufacture 2,5-dimethoxyamphetamine is consistent with the public interest at this time. DEA has investigated Pressure Chemical Company to ensure that the company's continued registration is consistent with the public interest. These investigations 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: March 27, 2002.

Laura M. Nagel,

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

[FR Doc. 02–8672 Filed 4–9–02; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated October 5, 2001, and published in the **Federal Register** on October 17, 2001, (66 FR 52782), Wildlife Laboratories, Inc., 1401 Duff Drive, Suite 600, Fort Collins, Colorado 80524, made application by renewal 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 firm plans to manufacture the listed controlled substance for distribution to its customers.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, section 823(a) and determined that the registration of Wildlife Laboratories to manufacture carfentanil is consistent with the public interest at this time. DEA has investigated Wildlife Laboratories to ensure that the company's registration is consistent with the public interest. This 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.