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]

BILLING CODE 4410–09–M

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.

Dated: March 27, 2002.

Laura M. Nagel,

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

[FR Doc. 02–8660 Filed 4–9–02; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Federal Bureau of Investigation

Meeting of the Compact Council for the National Crime Prevention and Privacy Compact

AGENCY: Federal Bureau of Investigation, Justice.

ACTION: Meeting notice.

SUMMARY: The purpose of this notice is to announce a meeting of the Compact Council created by the National Crime Prevention and Privacy Compact Act of 1998 (Compact). Thus far, the federal government and fourteen states are parties to the Compact which governs the exchange of criminal history records for licensing, employment, and similar purposes. The Compact also provides a legal framework for the establishment of a cooperative Federal-State system to exchange such records.

Matters for discussion are expected to include: (1) Dispute Adjudication Procedures, (2) Memorandum of Understanding with Nonparty States, (3) Expansion of the National Fingerprint File Participants, (4) Privatization of Noncriminal Justice Functions, and (5) Improvements to Background Checks and the use of Flat Fingerprints.

The meeting will be open to the public on a first-come, first-seated basis. Any member of the public wishing to file a written statement with the Compact Council or wishing to address this session of the Compact Council should notify Ms. Cathy L. Morrison at (304) 625–2736, at least 24 hours prior to the start of the session. The notification should contain the requestor's name and corporate designation, consumer affiliation, or government designation, along with a short statement describing the topic to be addressed, and the time needed for the presentation. Requestors will ordinarily be allowed up to 15 minutes to present a topic.

DATES AND TIMES: The Compact Council will meet in open session from 9 a.m. until 5 p.m. on May 8–9, 2002.

ADDRESSES: The meeting will take place at the Renaissance Scottsdale Resort, 6160 North Scottsdale Road, Scottsdale, Arizona, telephone (480) 991–1414.

FOR FURTHER INFORMATION CONTACT: Inquiries may be addressed to Ms. Cathy L. Morrison, Interim Compact Officer, Compact Council Office, Module C3, 1000 Custer Hollow Road, Clarksburg, West Virginia 26306–0147, telephone (304) 625–2736, facsimile (304) 625–5388.

Dated: March 21, 2002.

Thomas E. Bush, III,

Section Chief, Programs Development Section, Federal Bureau of Investigation.

[FR Doc. 02–8682 Filed 4–9–02; 8:45 am]

BILLING CODE 4410–02–M

NUCLEAR REGULATORY COMMISSION

Agency Information Collection Activities; Proposed Collection; Comment Request

AGENCY: U.S. Nuclear Regulatory Commission (NRC).

ACTION: Notice of the OMB review of information collection and solicitation of public comment.

SUMMARY: The NRC is preparing a submittal to OMB for review of continued approval of information collections under the provisions of Paperwork Reduction Act of 1995 (44 U.S.C. chapter 35).

Information pertaining to the requirement to be submitted:

1. *The title of the information collection:* 10 CFR Part 32—Specific Domestic Licenses to Manufacture or Transfer Certain Items Containing Byproduct Material.

2. *Current OMB approval number:* 3150–0001.

3. *How often the collection is required:* There is a one-time submittal of information to receive a license. Renewal applications are submitted every 10 years. In addition, recordkeeping must be performed on an on-going basis, and reports of transfer of byproduct material must be reported every 10 years.

4. *Who is required or asked to report:* All specific licensees who manufacture or initially transfer items containing byproduct material for sale or distribution to general licensees or persons exempt from licensing.

5. *The number of annual respondents:* 194 NRC licensees and 491 Agreement State licensees.

6. *The number of hours needed annually to complete the requirement or request:* 151,644 (53,012 hours for NRC licensees [4,507 reporting + 48,505 hours recordkeeping]) or an average of 273 hours per licensee and (98,632 hours for Agreement State licensees [3,210 hours reporting + 95,422 hours recordkeeping]) or 201 hours per Agreement State licensee.

7. *Abstract:* 10 CFR part 32 establishes requirements for specific licenses for the introduction of byproduct material into products or materials and transfer of the products or materials to general licensees or persons exempt from licensing. It also prescribes requirements governing holders of the specific licenses. Some of the requirements are for information which must be submitted in an application for a specific license, records which must be kept, reports which must be submitted, and information which must be forwarded to general licensees and persons exempt from licensing. In addition, 10 CFR part 32 prescribes requirements for the issuance of certificates of registration (concerning radiation safety information about a product) to manufacturers or initial transferors of sealed sources and devices. Submission or retention of the information is mandatory for persons subject to the 10 CFR part 32 requirements. The information is used by NRC to make licensing and other regulatory determinations concerning the use of radioactive byproduct material in products and devices.

Submit, by June 10, 2002, comments that address the following questions:

1. Is the proposed collection of information necessary for the NRC to properly perform its functions? Does the information have practical utility?

2. Is the burden estimate accurate?

3. Is there a way to enhance the quality, utility, and clarity of the information to be collected?

4. How can the burden of the information collection be minimized, including the use of automated collection techniques or other forms of information technology?

A copy of the draft supporting statement may be viewed free of charge at the NRC Public Document Room located at One White Flint North, 11555 Rockville Pike, Rockville, MD. OMB clearance requests are available at the NRC worldwide web site (http://www.nrc.gov/public-involve/doc-comment/omb/index.html). The document will be available on the NRC home page site for 60 days after the signature date of this notice.

Comments and questions about the information collection requirements may be directed to the NRC Clearance Officer, Brenda Jo. Shelton, U.S. Nuclear Regulatory Commission, T–6 E 6, Washington, DC 20555–0001, by telephone at (301) 415–7233, or by Internet electronic mail at *INFOCOLLECTS@NRC.GOV.*