contested, the reason(s) for contesting it, and the proposed amendment thereof. Persons filing such requests should make the envelope with the following legend "Privacy Act Amendment Request."

#### RECORD SOURCE CATEGORIES:

Basic information contained in this index is gathered from INS inspections, adjudications, reports of investigation, sworn statements, correspondence and memoranda, official reports, memoranda, and written and electronic referrals from other government agencies, including Federal, State, and local, information from foreign government agencies and international organizations.

## SYSTEMS EXEMPTED FROM CERTAIN PROVISIONS OF THE ACT:

The Attorney General has exempted this system from subsections (c)(3) and (4), (d), (e)(1), (2), and (3), (e)(4)(G) and (H), (e)(5) and (8), and (g), of the Privacy Act pursuant to 5 U.S.C. 552a(j)(2). In addition, the Attorney General has exempted this system from subsections (c)(3), (d) and (e)(1), (e)(4)(G) and (H) of the Privacy Act pursuant to 5 U.S.C. 552a(k)(2). The exemptions apply only to the extent that records in the system are subject to exemptions pursuant to 5 U.S.C. 552a (j)(2) and (k)(2). INS has published proposed implementing regulations in accordance with the requirements of 5 U.S.C. 553(b), (c), and (e) and these are published in today's Federal Register.

[FR Doc. 01–8285 Filed 4–3–01; 8:45 am] BILLING CODE 4410–10–M

## **DEPARTMENT OF JUSTICE**

## DRUG ENFORCEMENT ADMINISTRATION

## Manufacturer of Controlled Substances; Notice of Registration

By Notice dated September 28, 2000, and published in the **Federal Register** on October 13, 2000, (65 FR 60978), Irix Pharmaceuticals, Inc., 101 Technology Place, Florence, South Carolina 29501, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of methylphenidate (1724), a basic class of controlled substance listed in Schedule II.

The firm plans to manufacture methylphenidate for demonstration purposes and for dosage form development and stability studies.

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 Irix Pharmaceuticals, Inc. to manufacture methylphenidate is consistent with the public interest at this time. DEA has investigated the firm on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, 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 26, 2001.

## Laura M. Nagel,

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

[FR Doc. 01–8180 Filed 4–3–01; 8:45 am] **BILLING CODE 4410–09–M** 

### **DEPARTMENT OF JUSTICE**

## **Drug Enforcement Administration**

# Manufacturer of Controlled Substances; Notice of Registration

By Notice dated September 6, 2000, and published in the **Federal Register** on September 25, 2000, (65 FR 57622), Noramco Inc., 1400 Olympic Drive, Athens, Georgia 30601, 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 below:

Drug	Schedule
Amphetamine (1100)	       

The firm plans to support its other manufacturing facility with manufacturing and analytical testing.

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. to manufacture the listed controlled

substances is consistent with the public interest at this time. DEA has investigated Noramco, Inc. on a regular basis to ensure that the company's registration is consistent with the public interest. These investigations have 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 classes of controlled substances listed above is granted.

Dated: March 14, 2001.

### Laura M. Nagel,

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

[FR Doc. 01–8182 Filed 4–3–01; 8:45 am] BILLING CODE 4410–09–M

#### **DEPARTMENT OF JUSTICE**

## **Drug Enforcement Administration**

# Importation of Controlled Substances; Notice of Application

Pursuant to section 1008 of the Controlled Substances Import and Export Act (21 U.S.C. 958(i)), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in Schedule I or II and prior to issuing a regulation under Section 1002(a) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with section 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on November 16, 2000, Noramco of Delaware, Inc., 500 Old Swedes Landing Road, Wilmington, Delaware 19801, made application by renewal to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Opium, raw (9600)	II II

The firm plans to import the listed controlled substances for the bulk manufacture of other controlled substances.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of these basic classes of controlled substances may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.43 in such forms as prescribed by 21 CFR 1316 47

Any such comments, objections, or requests for a hearing may be addressed, in quintuplicate, to the Deputy Assistant 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 not later than May 4, 2001.

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1301.34(b), (c), (d), (e), and (f). As noted in a previous notice at 40 FR 43745-46 (September 23, 1975), all applicants for registration to import basic classes of any controlled substances in Schedule I or II are and will continue to be required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration that the requirements for such registration pursuant to 21 U.S.C. 958(a), 21 U.S.C. 823(a), and 21 CFR 1311.42(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: March 14, 2001.

## Laura M. Nagel,

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

[FR Doc. 01–8181 Filed 4–3–01; 8:45 am] BILLING CODE 4410–09-M

### **DEPARTMENT OF LABOR**

## **Employment and Training Administration**

## Youth Development Practitioner Apprenticeship Implementation Grants

**AGENCY:** Employment and Training Administration (ETA), Labor. **ACTION:** Notice of availability of funds and Solicitation for Grant Applications (SGAs).

This notice contains all of the necessary information and forms needed to apply for grant funding.

SUMMARY: The U.S. Department of Labor (DOL), Employment and Training Administration (ETA), announces competitive grants to be awarded under the Youth Development Practitioner

Apprenticeship initiative. This initiative targets incumbent and prospective professional youth workers working directly with young people. The funding available for these grants is \$1.45 million dollars and includes three distinct categories for application and award. The three categories are: (1) Funds for Local Intermediaries to Support Local Youth Program Service Operators in the Implementation of Apprenticeship Programs, (2) Grants to National Organizations, and (3) Provider of Technical Assistance on Practice and Curriculum Materials. An applicant can apply for more than one category of grant.

**DATES:** The closing date for receipt of applications is Friday, May 11, 2001. Applications must be received by 4:00 p.m. (Eastern Daylight Savings Time) at the address below. No exceptions to the mailing and hand-delivery conditions set forth in this notice will be granted. Applications that do not meet the conditions set forth in this notice will not be honored. Telefacsimile (FAX) applications will not be honored. **ADDRESSES:** Applications must be

mailed to: U.S. Department of Labor, Employment and Training Administration, Division of Federal Assistance, Attention: Mamie Williams, Reference: SGA/DFA 01–103, 200 Constitution Avenue, NW., Room S– 4203, Washington, DC 20210.

**Note:** Your application should specify on the cover which category you are applying for: 1, 2 or 3.

Hand Delivered Proposals. If proposals are hand delivered, they must be received at the designated address by 4:00 p.m., Eastern Daylight Time on Friday, May 11, 2001. All overnight mail will be considered to be hand delivered and must be received at the designated place by the specified closing date and time. Telegraphed, emailed and/or fax proposals will not be honored. Failure to adhere to the above instructions will be a basis for determination of non-responsive.

Late Proposals. A proposal received at the designated office after the exact time specified for receipt will not be considered unless it is received before the award is made and it:

• Was sent by U.S. Postal Service registered or certified mail not later than the fifth day (5th) calendar day before the closing date specified for receipt of applications (e.g. an offer submitted in response to a solicitation requiring receipt of applications by the 20th of the month must be mailed by the 15th):

• Was sent by U.S. Postal Service Express Mail Next Day Service, Post Office to Addressee, not later than 5 p.m. at the place of mailing two working days prior to the deadline date specified for receipt of proposals in this SGA. The term "working days" excludes weekends and U.S. Federal holidays.

The only acceptable evidence to establish the date of mailing of an application received after the deadline date for the receipt of proposals sent by the U.S. Postal Service and on the original receipt from the U.S. Postal Service. The term "post marked" means a printed, stamped, or otherwise placed impression (exclusive of a postage meter machine impression) that is readily identifiable without further action as having been supplied or affixed on the date of mailing by employees of the U.S. Postal Service.

Withdrawal of Applications.
Applications may be withdrawn by written notice or telegram (including mailgram) received at any time before an award is made. Applications may be withdrawn in person by the applicant or by an authorized representative thereof, if the representative's identity is made known and the representative signs a receipt for the proposal.

#### FOR FURTHER INFORMATION CONTACT:

Questions should be faxed to Mamie Williams at 202–693–2879, (this is not a toll-free number). All inquiries should include the SGA/DFA number 01–103, and a contact name, fax and phone numbers. This announcement will also be published on the Internet on the Employment and Training Administration's Home Page at http://www.doleta.gov. Award notifications will also be published on the Home Page.

## SUPPLEMENTARY INFORMATION:

## A. Authority

Section 171 of the Workforce Investment Act authorizes the use for demonstration program funds appropriated under section 174(b) for the purpose of developing and implementing techniques and approaches, and demonstrating the effectiveness of specialized methods, in addressing employment and training needs. Section 171(d) of the Workforce Investment Act authorizes the use for dislocated worker demonstration programs of funds reserved under section 132(a)(2)(A) and establishes the administration of these funds by the Secretary for that purpose under section 173(b). DOL FY 2000 Appropriations Act, enacted November 17, 1999, authorizes dislocated worker demonstration projects that provide assistance to new entrants in the workforce and incumbent workers. Apprenticeship programs are authorized