

firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: October 1, 1999.

**John H. King,**

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

[FR Doc. 99-26601 Filed 10-12-99; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Importation of Controlled Substances; Notice of Registration

By Notice dated July 1, 1999, and published in the **Federal Register** on August 2, 1999, (64 FR 41969), Calbiochem-Novabiochem Corporation, 10394 Pacific Center Court, Attn: Receiving Inspector, San Diego, California 92121-4340, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed below:

| Drug                               | Schedule |
|------------------------------------|----------|
| Tetrahydrocannabinols (7370) ..... | I        |
| Mescaline (7381) .....             | I        |
| Phencyclidine (7471) .....         | II       |
| Phenylacetone (8501) .....         | II       |
| Cocaine (9041) .....               | II       |

The firm plans to import small quantities of the listed controlled substances to make reagents for distribution to the biomedical research community.

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 Calbiochem-Novabiochem Corporation 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 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 Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title

21, Code of Federal Regulations, Section 1301.34, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: October 1, 1999.

**John H. King,**

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

[FR Doc. 99-26602 Filed 10-12-99; 8:45 am]

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## 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 August 16, 1999, Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Building 18, Chattanooga, Tennessee 37409, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of methamphetamine (1105), a basic class of controlled substance listed in Schedule II.

The firm plans to bulk manufacture methamphetamine to produce products for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance 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 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 no later than December 13, 1999.

Dated: October 1, 1999.

**John H. King,**

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

[FR Doc. 99-26604 Filed 10-12-99; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances; Notice of Registration

By Notice dated June 23, 1999, and published in the **Federal Register** on July 7, 1999, (64 FR 36717),

Damocles10, 3529 Lincoln Highway, Thorndale, Pennsylvania 19372, 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 |
|-------------------------------|----------|
| Codeine-N-oxide (9053) .....  | I        |
| Heroin (9200) .....           | I        |
| Morphine-N-oxide (9307) ..... | I        |
| Amphetamine (1100) .....      | II       |
| Methamphetamine (1105) .....  | II       |
| Phencyclidine (7471) .....    | II       |
| Codeine (9050) .....          | II       |
| Morphine (9300) .....         | II       |

The firm plans to manufacture the listed controlled substances for the purpose of deuterium labeled internal standards for distribution to analytical laboratories.

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 Damocles10 to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Damocles10 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 classes of controlled substances listed above is granted.

Dated: October 1, 1999.

**John H. King,**

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

[FR Doc. 99-26603 Filed 10-12-99; 8:45 am]

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## 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 (12 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 August 19, 1999, Fort Dodge Laboratories, Inc., 141 E. Riverside Drive, Fort Dodge, Iowa 50501, made application by renewal to the Drug Enforcement Administration to be registered as an importer of pentobarbital (2270), a basic class of controlled substance listed in Schedule II.

The firm plans to manufacture a product for distribution to its customers.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substance 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 form 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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than (30 days from publication).

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 class of any controlled substance 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: October 1, 1999.

**John H. King,**

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

[FR Doc. 99-26605 Filed 10-12-99; 8:45 am]

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## 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 August 9, 1999, Irix Pharmaceuticals, Inc., 101 Technology Place, Florence, South Carolina 29501, made application by renewal to the Drug Enforcement Administration (DEA) for registration 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.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance 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 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 no later than December 13, 1999.

Dated: October 1, 1999.

**John H. King,**

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

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## DEPARTMENT OF JUSTICE

### National Institute of Corrections

#### Solicitation for a Cooperative Agreement

**SUMMARY:** The Department of Justice (DOJ), National Institute of Corrections (NIC), announces the availability of funds in FY 2000 for a cooperative agreement to develop the training curriculum, How to Develop Management Training.

The National Institute of Corrections (NIC) invites applications for a cooperative agreement to develop a standard, core curriculum for training persons responsible for the development of management training for supervisors and administrators within juvenile corrections and detention settings. To enable the Institute to offer state-of-the-art guidance for the development of management training, the award recipient will develop a 32-hour training curriculum including an instructors' guide with lesson plans, computer-generated view graphs to support the curriculum, and participant manual. The 32-hour curriculum will provide juvenile corrections and detention trainers multiple development and delivery methods and strategies to construct management training within their agencies that will equip managers with the core competencies to perform effectively. (It is not within the scope of this cooperative agreement to provide piloting or direct delivery of the curriculum.)

The award recipient will become familiar with the management and leadership training programs currently being offered at NIC. The recipient will utilize this information, as well as contribute to the development of new information on management practices most desirable in today's rapidly changing juvenile corrections and detention environment.

As a collaborative venture with the NIC Academy Division, the recipient will develop training outcomes for the project in partnership with the NIC project manager. Funding for this cooperative agreement comes from an Interagency Agreement (IAA) between the Office of Juvenile Justice and Delinquency Prevention (OJJDP) and NIC. A total of \$30,000 is reserved for the project which will support one cooperative agreement for a 6-month period. The recipient of the award will be selected through a competitive solicitation process. Steven Swisher is the designated NIC project manager.

#### Background

Well-trained, effective managers and leaders within juvenile corrections and detention agencies have been a focus of the training and services the NIC has provided through an IAA with OJJDP over the past nine years. As a part of that IAA and as a result of a national juvenile training needs assessment conducted in the fall of 1998, curricula and services for training staff continue to be identified as critical in capacitating juvenile correctional and detention agencies to develop and sustain effective management and