proposed Consent Decree may be obtained in person or by mail from the Consent Decree Library, 1120 G Street, NW., 4th Floor, Washington, DC 20005. In requesting a copy please refer to the referenced case and enclose a check in the amount of $4.25 (25 cents per page reproduction costs), payable to the Consent Decree Library.

**Joel M. Gross,**

*Chief, Environmental Enforcement Section, Environment and Natural Resources Division.*

[FR Doc. 98–18253 Filed 7–8–98; 8:45 am]

**BILLING CODE 4410–15–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 April 1, 1998, Aernol Pharmaceutical, Inc., 189 Meister Avenue, Somerville, New Jersey 08876, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below.

| Drug | Schedule |
|---|---|
| N-Ethylamphetamine (1475) | I |
| Difenoxin (9168) | I |
| Amphetamine (1100) | II |
| Methamphetamine (1105) | II |

The firm plans to manufacture the listed controlled substances to produce pharmaceutical products for its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture 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 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 September 8, 1998.

Dated: June 10, 1998.

**John H. King,**

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

[FR Doc. 98–18217 Filed 7–8–98; 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 January 28, 1998, Chiragene, Inc., 7 Powder Horn Drive, Warren, New Jersey 07059, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below.

| Drug | Schedule |
|---|---|
| N-Ethylamphetamine (1475) | I |
| 2,5-Dimethoxyamphetamine (7396). | I |
| 3,4-Methylenedioxyamphetamine (7400). | I |
| 4-Methoxyamphetamine (7411) | I |
| Amphetamine (1100) | II |
| Methylphenidate (1724) | II |

The firm plans to manufacture the listed controlled substances to supply their customers.

Any other such applicant and any person who is presently registered with DEA to manufacture 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 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 September 8, 1998.

Dated: June 10, 1998.

**John H. King,**

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

[FR Doc. 98–18216 Filed 7–8–98; 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 April 19, 1998, Damocles 10, 3529 Lincoln Highway, Thorndale, Pennsylvania 19372, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below.

| Drug | Schedule |
|---|---|
| Heroin (9200) | I |
| Amphetamine (1100) | II |
| Methamphetamine (1105) | II |
| Phenmetrazine (1631) | II |
| Hydromorphone (9150) | 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.

Any other such applicant and any person who is presently registered with DEA to manufacture 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 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 September 8, 1998.

Dated: June 30, 1998.

**John H. King,**

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

[FR Doc. 98–18219 Filed 7–8–98; 8:45 am]

**BILLING CODE 4410–09–M**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Registration

By notice dated January 21, 1998, and published in the **Federal Register** on February 12, 1998 (63 FR 7181), Johnson & Johnson Pharmaceutical Partners, HC–02 Box 19250, KMO.1 Mamey Ward (HC–02 Box 19250), Gurabo, Puerto Rico 00778–9629, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of sufentanil (9740), a basic class of controlled substance listed in schedule II.

The firm plans to manufacture sufentanil for bulk distribution to its customers.

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Johnson & Johnson Pharmaceutical Partners to manufacture sufentanil is consistent with the public interest at this time. Therefore, pursuant