The proposed Consent Decree may be examined at the office of the United States Attorney, Federico Degetau Federal Bldg., Carlos E. Chardon Avenue, Hato Rey, Puerto Rico 00918 and at the Region II office of the Environmental Protection Agency, 290 Broadway, New York, New York 10007. The proposed Consent Decree may also be examined at the Consent Decree Library, 1120 G St., NW., 4th Floor, Washington, DC 20005, 202–624–0892. A copy of the proposed Consent Decree may be obtained in person or by mail from the Consent Decree Library, 1120 G St., NW., 4th Floor, Washington, DC 20005. In requesting a copy, please enclose a check in the amount of $6.50 (25 cents per page reproduction cost) payable to the ''Consent Decree Library.''

**Walker B. Smith,**

*Deputy Chief, Environmental Enforcement Section, Environment and Natural Resource Division.*

[FR Doc. 97–11947 Filed 5–7–97; 8:45 am]

**BILLING CODE 4410–15–M**

## DEPARTMENT OF JUSTICE

### Antitrust Division

### U.S. v. Restonic Corporation, Serta, Incorporated, and Spring Air Company

Notice is hereby given that defendants Restonic Corporation (''Restonic''); Serta, Inc. (formerly Serta Associates, Inc.) (''Serta''); and Spring Air Company (''Spring Air'') have filed with the United States District Court for the Northern District of Illinois motions to terminate the Judgments in *United States* v. *Restonic Corporation,* Civil No. 60–C–828; *United States* v. *Serta, Inc.,* Civil No. 60–C–843; and *United States* v. *Spring Air Company,* Civil No. 60–C–845. In Stipulations also filed with the Court, the Department of Justice (''Department'') has tentatively consented to termination of the Judgments, but has reserved the right to withdraw its consent pending receipt of public comments.

The complaints in these cases, filed on May 27, 1960 in the case of Restonic and on May 31, 1960 with respect to the other defendants, alleged that each defendant had conspired with its respective licensee owners to illegally allocate exclusive territories and to fix retail prices. Judgments were entered by consent against Restonic on May 27, 1960 (and subsequently modified on July 27, 1962); and against Spring Air on May 31, 1960. Judgment was entered after a trial on the merits against Serta on September 30, 1968. The Judgments prohibited defendants from establishing territories for the sale of mattresses and from engaging in resale price maintenance.

The Department has filed with the Court a memorandum setting forth the reasons why the Government believes that termination of the Judgments would serve the public interest. Copies of the Defendants' motion papers, the Stipulations containing the Government's consent, the Government's memorandum and all further papers filed with the court in connection with this motion will be available for inspection at the Legal Procedure Unit of the Antitrust Division, Room 215 North, Liberty Place Building, 325 Seventh St., N.W., Washington, D.C. 20530, and at the Office of the Clerk of the United States District Court for the Northern District of Illinois, Twentieth Floor, 209 South Dearborn Street, Chicago, Illinois 60604. Copies of these materials may be obtained from the Antitrust Division upon request and payment of the copying fee set by Department of Justice regulations.

Interested persons may submit comments regarding the proposed termination of the Judgments to the Department. Such comments must be received by the Department within sixty (60) days and will be filed with the Court by the Department. Comments should be addressed to Mildred L. Calhoun, Trail Attorney, Midwest Office, Antitrust Division, Suite 600, 209 S. LaSalle Street, Chicago, Illinois 60604 (Telephone: (312) 353–7530.

**Rebecca P. Dick,**

*Deputy Director of Operations.*

[FR Doc. 97–11940 Filed 5–7–97; 8:45 am]

**BILLING CODE 4410–11–M**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Registration

By Notice dated December 31, 1996, and published in the **Federal Register** on January 9, 1997 (62 FR 1342), Ansys, Inc., 2 Goodyear, Irvine, California 92718, 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 |
|---|---|
| Phencyclidine (7471) | II |
| 1-Piperidinocyclohexanecarbonitrile (PCC) (8603). | II |
| Benzoylecgonine (9180) | II |

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 Ansys, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 0.100 and 0.104, the Acting 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: April 21, 1997.

**Terrance W. Woodworth,**

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

[FR Doc. 97–11916 Filed 5–7–97; 8:45 am]

**BILLING CODE 4410–09–M**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Registration

By Notice dated December 18, 1996, and published in the **Federal Register** on January 9, 1997 (62 FR 1342), Arenol Chemical Corporation, 189 Meister Avenue, Somerville, New Jersey 08876, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of N-ethylamphetamine (1475), a basic class of controlled substance listed in Schedule I.

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 Arenol Chemical Corporation to manufacture n-ethylamphetamine is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 0.100 and 0.104, the Acting 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: April 4, 1997.

**Terrance W. Woodworth,**

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

[FR Doc. 97–11917 Filed 5–7–97; 8:45 am]

BILLING CODE 4410–09–M

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.43(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on February 26, 1997, Ganes Chemicals, Inc., Industrial Park Road, Pennsville, New Jersey 08070, 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 |
|---|---|
| Methylphenidate (1724) | II |
| Amobarbital (2125) | II |
| Pentobarbital (2270) | II |
| Secobarbital (2315) | II |
| Glutethimide (2550) | II |
| Methadone (9250) | II |
| Methadone-intermediate (9254) | II |
| Dextropropoxyphene, bulk (non-dosage forms) (9273). | II |

The firm plans to manufacture the controlled substances for distribution as bulk products to 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 above application.

Any such comments or objections may be addressed, in quintuplicate, to the Acting 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 July 7, 1997.

Dated: March 31, 1997.

**Terrance W. Woodworth,**

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

[FR Doc. 97–11918 Filed 5–7–97; 8:45 am]

BILLING CODE 4410–09–M

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Correction

As set forth in the **Federal Register** (FR Doc. 96–32611) Vol. 61, No. 248 at page 67852, dated December 24, 1996, High Standard Products, 1100 W. Florence Avenue, #B, Inglewood, California 90301, made application to the Drug Enforcement Administration for registration as a manufacturer for certain controlled substances.

By letter dated March 19, 1997, High Standard Products stated that they had erroneously included marihuana (7360) in their application. Therefore, marihuana is hereby deleted from the firm's application for registration as a manufacturer.

Dated: March 31, 1997.

**Terrance W. Woodworth,**

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

[FR Doc. 97–11919 Filed 5–7–97; 8:45 am]

BILLING CODE 4410–09–M

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Registration

By Notice dated November 18, 1996, and published in the **Federal Register** on December 24, 1996, (61 FR 67852), High Standard Products, 1100 W. Florence Avenue, #B, Inglewood, California 90301, made application by renewal to the Drug Enforcement Administration (DEA) to be registered to manufacture small quantities of controlled substances for drug reference standards as a bulk manufacturer of the basic classes of controlled substances listed below:

| Drug | Schedule |
|---|---|
| Methaqualone (2565) | I |
| Lysergic acid diethylamide (7315) | I |
| Tetrahydrocannabinols (7370) | I |
| 3,4-Methylenedioxyamphetamine (7400). | I |
| 3,4-Methylenedioxy-N-ethylamphetamine (7404). | I |
| 3,4-Methylenedioxymethamphetamine (7405). | I |
| 4-Methoxyamphetamine (7411) | I |
| 1-(1-Phenylcyclohexyl)pyrrolidine (PCPY) (7458). | I |
| Heroin (9200) | I |
| Normorphine (9313) | I |
| 3-Methylfentanyl (9813) | I |
| Amphetamine (1100) | II |
| Methamphetamine (1105) | II |
| Secobarbital (2315) | II |
| Phencyclidine (7471) | II |
| Cocaine (9041) | II |
| Codeine (9050) | II |
| Hydromorphone (9150) | II |
| Diphenoxylate (9170) | II |
| Benzoylecgonine (9180) | II |
| Hydrocodone (9193) | II |
| Methadone (9250) | II |
| Morphine (9300) | II |
| Fentanyl (9801) | II |

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 High Standard Products to manufacture the listed controlled substances is consistent with the public interest at this time. Therefore, pursuant to 21 USC § 823 and 28 CFR §§ 0.100 and 0.104, the Acting Deputy Assistant Administration, Office of Division 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: April 16, 1997.

**Terrance W. Woodworth,**

*Acting Deputy Assistant Administrator, Drug Enforcement Administration.*

[FR Doc. 97–11920 Filed 5–7–97; 8:45 am]

BILLING CODE 4410–09–M

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.43(a) of Title 21 of the code of Federal Regulations (CFR), this is notice that on December 18, 1996, Johnson Matthey, Inc., Custom Pharmaceuticals Department, 2003 Nolte Drive, West Deptford, NJ 08066, 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 |
|---|---|
| Difenoxin (9168) | I |
| Methylphenidate (1724) | II |
| Dihydrocodeine (9120) | II |
| Oxycodone (9143) | II |
| Hydromorphone (9150) | II |
| Hydrocodone (9193) | II |
| Levorphanol (9220) | II |
| Meperidine (9230) | II |
| Methadone (9250) | II |
| Thebaine (9333) | II |