

Portland, OR; VicTrack, Docklands, AUSTRALIA; VIP Operator, Skopje, MACEDONIA; Virtus IT Limited, London, UNITED KINGDOM; VISITEK, Jakarta Selatan, INDONESIA; Vitria Technology, Inc., Sunnyvale, CA; Vonage, Holmdel, NJ; Wataniya Telecom Kuwait, Plot 1A, Sharq Area, KUWAIT; Wind Telecomunicazioni SpA, Roma, ITALY; WiTech, Cascina, ITALY; Xelas software, Marina del Rey, CA; XTRAC, LLC, Boston, MA; Yyield Group BV, Bennebroek, NETHERLANDS; Zain, Safat, KUWAIT; ZAO "Glasnet", Korolev, RUSSIA; Zenoss, Annapolis, MD and Zenulta Limited, Swindon, UNITED KINGDOM.

No other changes have been made in either the membership or planned activity of the group research project. Membership in this group research project remains open, and The Forum intends to file additional written notifications disclosing all changes in membership.

On October 21, 1988, The Forum filed its original notification pursuant to Section 6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to Section 6(b) of the Act on December 8, 1988 (53 FR 49615).

The last notification was filed with the Department on August 22, 2011. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on November 18, 2011 (76 FR 71602).

**Patricia A. Brink,**

*Director of Civil Enforcement, Antitrust Division.*

[FR Doc. 2012-30724 Filed 12-20-12; 8:45 am]

**BILLING CODE P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Importer of Controlled Substances, Notice of Application, Hospira

Pursuant to Title 21 Code of Federal Regulations 1301.34(a), this is notice that on September 20, 2012, Hospira, 1776 North Centennial Drive, McPherson, Kansas 67460-1247, made application by renewal to the Drug Enforcement Administration (DEA) for registration as an importer of Remifentanyl (9739), a basic class of controlled substance listed in schedule II.

The company plans to import Remifentanyl for use in dosage form manufacturing.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture

such basic class of controlled substance may file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43, and in such form as prescribed by 21 CFR 1316.47.

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, **Federal Register** Representative (ODL), 8701 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than January 22, 2013.

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 published in the **Federal Register** on September 23, 1975, 40 FR 43745-46, all applicants for registration to import a basic class of any controlled substance in schedules 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 1301.34(b), (c), (d), (e), and (f) are satisfied.

Dated: December 14, 2012.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2012-30782 Filed 12-20-12; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Importer of Controlled Substances; Notice of Registration; Fisher Clinical Services, Inc.

By Notice dated September 20, 2012, and published in the **Federal Register** on October 2, 2012, 77 FR 60143, Fisher Clinical Services, Inc., 7554 Schantz Road, Allentown, Pennsylvania 18106, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of noroxymorphone (9668), a basic class of controlled substance in schedule II.

The company plans to import the listed substances for analytical research and clinical trials.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and 952(a) and determined that the registration of Fisher Clinical Services, Inc., to import

the basic class of controlled substance is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. DEA has investigated Fisher Clinical Services, Inc., to ensure that the company's registration is consistent with the public interest. The investigation has 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. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above named company is granted registration as an importer of the basic class of controlled substance listed.

Dated: December 14, 2012.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2012-30784 Filed 12-20-12; 8:45 am]

**BILLING CODE 4410-09-P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances, Notice of Registration, AMRI Rensselaer, Inc.

By Notice dated July 30, 2012, and published in the **Federal Register** on August 7, 2012, 77 FR 47114, AMRI Rensselaer, Inc., 33 Riverside Avenue, Rensselaer, New York 12144, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

| Drug                                            | Schedule |
|-------------------------------------------------|----------|
| Marihuana (7360) .....                          | I        |
| Tetrahydrocannabinols (7370) .....              | I        |
| Amphetamine (1100) .....                        | II       |
| Lisdexamfetamine (1205) .....                   | II       |
| Methylphenidate (1724) .....                    | II       |
| Pentobarbital (2270) .....                      | II       |
| 4-Anilino-N-phenethyl-4-piperidine (8333) ..... | II       |
| Meperidine (9230) .....                         | II       |
| Fentanyl (9801) .....                           | II       |

The company plans to manufacture bulk controlled substances for use in product development and for distribution to its customers.

In reference to drug code 7360 (Marihuana), the company plans to bulk manufacture cannabidiol as a synthetic intermediate. This controlled substance

will be further synthesized to bulk manufacture a synthetic THC (7370). No other activity for this drug code is authorized for this registration.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a), and determined that the registration of AMRI Rensselaer, Inc., to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated AMRI Rensselaer, Inc., to ensure that the company's registration is consistent with the public interest. The investigation has 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 in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: December 18, 2012.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2012-30780 Filed 12-20-12; 8:45 am]

**BILLING CODE 4410-09-P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

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

By Notice dated May 31, 2012, and published in the **Federal Register** on June 8, 2012, 77 FR 34072, Rhodes Technologies, 498 Washington Street, Coventry, Rhode Island 02816, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of morphine (9300), a basic class of controlled substance listed in schedule II.

The company plans to manufacture the listed controlled substance in bulk for conversion and sale to dosage form manufacturers.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a), and determined that the registration of Rhodes Technologies to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Rhodes Technologies to ensure that the company's registration is consistent with the public interest. The

investigation has 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(a), and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic class of controlled substance listed.

Dated: December 14, 2012.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2012-30785 Filed 12-20-12; 8:45 am]

**BILLING CODE 4410-09-P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances; Notice of Registration; Euticals, Inc.

By Notice dated August 17, 2012, and published in the **Federal Register** on August 29, 2012, 77 FR 52367, Euticals, Inc., (formerly known as Archimica, Inc.), 2460 W. Bennett Street, Springfield, Missouri 65807-1229, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

| Drug                              | Schedule |
|-----------------------------------|----------|
| Gamma Hydroxybutyric Acid (2010). | I        |
| Amphetamine (1100) .....          | II       |
| Lisdexamfetamine (1205) .....     | II       |
| Methylphenidate (1724) .....      | II       |
| Phenylacetone (8501) .....        | II       |
| Methadone Intermediate (9254) ... | II       |
| Tapentadol (9780) .....           | II       |

The company plans to manufacture the listed controlled substances in bulk for distribution and sale to its customers.

With regards to amphetamine (1100), the company plans to acquire the listed controlled substance in bulk from a domestic source in order to manufacture other controlled substances in bulk for distribution to its customers.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a), and determined that the registration of Euticals, Inc., to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Euticals,

Inc., to ensure that the company's registration is consistent with the public interest. The investigation has 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(a), and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: December 14, 2012.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2012-30783 Filed 12-20-12; 8:45 am]

**BILLING CODE 4410-09-P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances; Notice of Registration; Chemtos, LLC

By Notice dated July 30, 2012, and published in the **Federal Register** on August 7, 2012, 77 FR 47116, Chemtos, LLC, 14101 W. Highway 290, Building 2000B, Austin, Texas 78737-9331, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

| Drug                                | Schedule |
|-------------------------------------|----------|
| Amphetamine (1100) .....            | II       |
| Methamphetamine (1105) .....        | II       |
| Lisdexamfetamine (1205) .....       | II       |
| Methylphenidate (1724) .....        | II       |
| Nabilone (7379) .....               | II       |
| Phenylacetone (8501) .....          | II       |
| Cocaine (9041) .....                | II       |
| Codeine (9050) .....                | II       |
| Etorphine HCL (9059) .....          | II       |
| Dihydrocodeine (9120) .....         | II       |
| Oxycodone (9143) .....              | II       |
| Hydromorphone (9150) .....          | II       |
| Ecgonine (9180) .....               | II       |
| Ethylmorphine (9190) .....          | II       |
| Hydrocodone (9193) .....            | II       |
| Levomethorphan (9210) .....         | II       |
| Levorphanol (9220) .....            | II       |
| Isomethadone (9226) .....           | II       |
| Meperidine (9230) .....             | II       |
| Meperidine-intermediate-A (9232) .. | II       |
| Meperidine-intermediate-B (9233) .. | II       |
| Meperidine-intermediate-C (9234) .. | II       |
| Methadone (9250) .....              | II       |
| Methadone intermediate (9254) ...   | II       |
| Morphine (9300) .....               | II       |
| Thebaine (9333) .....               | II       |
| Dihydroetorphine (9334) .....       | II       |
| Levo-alphaacetylmethadol (9648) ..  | II       |