

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.*

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**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.*

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**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.*

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**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