

Dated: January 6, 2012.

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

[FR Doc. 2012-679 Filed 1-13-12; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled  
Substances; Notice of Application**

Pursuant to § 1301.33(a), of Title 21 of the Code of Federal Regulations (CFR), this is notice that on September 6, 2011 Cedarburg Pharmaceuticals, Inc., 870 Badger Circle, Grafton, Wisconsin 53024, 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
4-Anilino-N-phenethyl-4-piperidine (8333).	II
Fentanyl (9801) .....	II

The company plans to manufacture the listed controlled substances in bulk for distribution to its customers. Regarding the drug code (8333), the company plans to use this controlled substance to manufacture another controlled substance.

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 pursuant to 21 CFR 1301.33(a).

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 March 19, 2012.

Dated: January 6, 2012.

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

[FR Doc. 2012-656 Filed 1-13-12; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled  
Substances; Notice of Application**

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on September 12, 2011, Johnson Matthey, Inc., Pharmaceuticals Materials, 900 River Road, Conshohocken, Pennsylvania 19428, 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
Methylphenidate (1724) .....	II
Codeine (9050) .....	II
Oxycodone (9143) .....	II
Hydrocodone (9193) .....	II
Morphine (9300) .....	II
Thebaine (9333) .....	II

The company plans to manufacture the listed controlled substances in bulk for sale in bulk to its customers. The Thebaine (9333) will also be used to manufacture other controlled substances in bulk which will also be for sale in bulk 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 proposed registration pursuant to 21 CFR 1301.33(a).

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 March 19, 2012.

Dated: January 6, 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**

By Notice dated August 9, 2011, and published in the **Federal Register** on August 18, 2011, 76 FR 51401, Austin

Pharma LLC., 811 Paloma Drive, Suite C, Round Rock, Texas 78665-2402, 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

The company plans to manufacture bulk active pharmaceutical ingredients (APIs) 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 Austin Pharma LLC. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Austin Pharma LLC. 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: January 6, 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**

By Notice dated August 10, 2011, and published in the **Federal Register** on August 18, 2011, 76 FR 51401, AMRI Rensselaer, Inc., 33 Riverside Avenue, Rensselaer, New York 12144, made application by renewal to the Drug