

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled Substances; Notice of Application; Chemtos, LLC.**

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on June 14, 2012, 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
Oxymorphone (9652) .....	II
Racemethorphan (9732) .....	II
Racemorphan (9733) .....	II

The company plans to manufacture small quantities of the listed controlled substances in bulk for distribution to its customers for use as reference standards.

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 October 9, 2012.

Dated: July 30, 2012.

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

[FR Doc. 2012-19187 Filed 8-6-12; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled Substances; Notice of Registration; Penick Corporation**

By Notice dated April 17, 2012, and published in the **Federal Register** on April 26, 2012, 77 FR 24986, Penick Corporation, 33 Industrial Park Road, Pennsville, New Jersey 08070, 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
Cocaine (9041) .....	II
Codeine (9050) .....	II
Dihydrocodeine (9120) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Ecgonine (9180) .....	II
Hydrocodone (9193) .....	VII
Morphine (9300) .....	II
Oripavine (9330) .....	II
Thebaine (9333) .....	II
Oxymorphone (9652) .....	II

The company plans to manufacture the listed controlled substances as bulk controlled substance intermediates 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 Penick Corporation to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time.

DEA has investigated Penick Corporation 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: July 30, 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; Cambrex Charles City, Inc.**

By Notice dated April 17, 2012, and published in the **Federal Register** on April 26, 2012, 77 FR 24986, Cambrex Charles City, Inc., 1205 11th Street, Charles City, Iowa 50616, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
Cocaine (9041) .....	II
Opium tincture (9630) .....	II

The company plans to manufacture the listed controlled substances in bulk for sale 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 Cambrex Charles City, Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time.

DEA has investigated Cambrex Charles City, 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: July 30, 2012.

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

[FR Doc. 2012-19180 Filed 8-6-12; 8:45 am]

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