Dated: February 19, 2014.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2014–05496 Filed 3–12–14; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Registration; Morton Grove Pharmaceuticals

By Notice dated November 5, 2013, and published in the **Federal Register** on November 18, 2013, 78 FR 69133, Morton Grove Pharmaceuticals, 6451 Main Street, Morton Grove, Illinois 60053–2633, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Gamma Hydroxybutyric Acid (2010), a basic class of controlled substance listed in schedule I.

The company plans to manufacture a controlled substance for product development.

No comments or objections have been received. The DEA has considered the factors in 21 U.S.C. 823(a), and determined that the registration of Morton Grove Pharmaceuticals to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. The DEA has investigated Morton Grove Pharmaceuticals 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: February 19, 2014.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2014–05505 Filed 3–12–14; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Registration; Stepan Company

By Notice dated October 9, 2013, and published in the **Federal Register** on October 25, 2013, 78 FR 64018, Stepan Company, Natural Products Dept., 100 W. Hunter Avenue, Maywood, New Jersey 07607, 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 |
| Ecgonine (9180) | II |

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

No comments or objections have been received. The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Stepan Company to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. The DEA has investigated Stepan Company 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: February 19, 2014.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2014–05494 Filed 3–12–14; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Registration; Research Triangle Institute

By Notice dated October 10, 2013, and published in the **Federal Register** on October 25, 2013, 78 FR 64018, Research Triangle Institute, Hermann Building East Institute Drive, P.O. Box 12194, Research Triangle Park, North Carolina 27709, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Tetrahydrocannabinols (7370), a basic class of controlled substance listed in schedule I.

The company plans to provide small quantities to commercial customers for use in preparing test kits, reagents, and reference standards.

The company plans to bulk manufacture a synthetic Tetrahydrocannabinol. No other activity for this drug code is authorized for this registration.

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

Dated: February 19, 2014.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2014–05507 Filed 3–12–14; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Registration; Nektar Therapeutics

By Notice dated September 27, 2013, and published in the **Federal Register** on October 25, 2013, 78 FR 64018, Nektar Therapeutics, 1112 Church Street, Huntsville, Alabama 35801, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Fentanyl (9801), a basic

class of controlled substance listed in schedule II.

The company plans to manufacture the listed controlled substance in support of product development.

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

Dated: Signed February 19, 2014.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2014–05489 Filed 3–12–14; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Registration; Johnson Matthey, Inc.

By Notice dated November 5, 2013, and published in the **Federal Register** on November 18, 2013, 78 FR 69133, 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 |
| Diphenoxylate (9170) | II |
| Hydrocodone (9193) | II |
| Meperidine (9230) | II |
| Methadone (9250) | II |
| Methadone intermediate (9254) | II |
| Morphine (9300) | II |
| Thebaine (9333) | II |

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

No comments or objections have been received. The DEA has considered the factors in 21 U.S.C. 823(a), and determined that the registration of Johnson Matthey, Inc., to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. The DEA has investigated, Johnson Matthey, 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: February 19, 2014.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2014–05500 Filed 3–12–14; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

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

By Notice dated September 27, 2013, and published in the **Federal Register** on October 25, 2013, 78 FR 64017, PharmaCore, Inc., 4180 Mendenhall Oaks Parkway, High Point, North Carolina 27265, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Noroxymorphone (9668), a basic class of controlled substance listed in schedule II.

The company plans to manufacture the listed controlled substance as active pharmaceutical ingredients (API) for clinical trials.

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

Dated: February 19, 2014.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2014–05504 Filed 3–12–14; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Registration; Cambrex Charles City, Inc.

By Notice dated September 27, 2013, and published in the **Federal Register** on October 25, 2013, 78 FR 64017, Cambrex Charles City, Inc., 1205 11th Street, Charles City, Iowa 50616, 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 |
| 4-Anilino-N-phenethyl-4-piperidine (8333). | II |
| Phenylacetone (8501) | II |
| Cocaine (9041) | II |
| Codeine (9050) | II |
| Oxycodone (9143) | II |
| Hydromorphone (9150) | II |
| Hydrocodone (9193) | II |
| Methadone (9250) | II |
| Morphine (9300) | II |
| Oripavine (9330) | II |
| Thebaine (9333) | II |
| Opium, raw (9600) | II |
| Opium extracts (9610) | II |
| Opium fluid extract (9620) | II |
| Opium tincture (9630) | II |
| Opium, powdered (9639) | II |
| Opium, granulated (9640) | II |
| Oxymorphone (9652) | II |