The company plans to manufacture the listed controlled substances in bulk for sales to 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, 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: September 20, 2006.

**Joseph T. Rannazzisi,**

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

[FR Doc. E6–16030 Filed 9–28–06; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 18, 2006, Cambridge Isotope Lab, 50 Frontage Road, Andover, Massachusetts 01810, made application by renewal 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 utilize small quantities of the listed controlled substance in the preparation of analytical standards.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance 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 being sent via regular mail should be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA **Federal Register** Representative/ODL; or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA **Federal Register** Representative/ ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than November 28, 2006.

September 20, 2006.

**Joseph T. Rannazzisi,**

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

[FR Doc. E6–16029 Filed 9–28–06; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Importer of Controlled Substances; Notice of Application

Pursuant to 21 U.S.C. 958(i), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in Schedule I or II and prior to issuing a regulation under 21 U.S.C. 952(a)(2)(B) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with 21 CFR 1301.34(a), this is notice that on February 10, 2006, Dakota Pharmaceutical Packaging, 4733 Amber Valley Parkway, Fargo, North Dakota, 58104, made application to the Drug Enforcement Administration (DEA) to be registered as an Importer of Hydrocodone (9193), a basic class of controlled substances in Schedule II.

The company plans to import the listed controlled substance for clinical trials.

Any manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances 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 being sent via regular mail should be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, *Attention:* DEA **Federal Register** Representative/ODL; or any being sent via express mail should be sent to DEA Headquarters, *Attention:* DEA **Federal Register** Representative/ ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than October 30, 2006.

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 listed in Schedule 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 USC 958(a), 21 U.S.C. 823(a), and 21 CFR 1301.34(b), (c), (d), (e), and (f) are satisfied.

Dated: September 20, 2006.

**Joseph T. Rannazzisi,**

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

[FR Doc. E6–16053 Filed 9–28–06; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Registration

By Notice dated May 9, 2006, and published in the **Federal Register** on May 15, 2006, (71 FR 28052), Johnson Matthey Inc., Custom Pharmaceuticals Department, 2003 Nolte Drive, West Deptford, New Jersey 08066, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in Schedule II:

| Drug | Schedule |
|---|---|
| Dihydrocodeine (9120) | II |
| Oxymorphone (9652) | II |

The company plans to manufacture in bulk for distribution to its customers, who are final dosage 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 Johnson Matthey Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. 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, 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: September 20, 2006.

**Joseph T. Rannazzisi,**

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

[FR Doc. E6–16023 Filed 9–28–06; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 30, 2006, Abbott Laboratories, DBA Knoll Pharmaceutical Company, 30 North Jefferson Road, Whippany, New Jersey, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in Schedules I and II:

| Drug | Schedule |
|---|---|
| Dihydromorphine (9145) | I |
| Hydromorphone (9150) | II |

The company plans to manufacture bulk product and dosage units for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance 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 being sent via regular mail should be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative/ODL; or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA Federal Register Representative/ ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than November 28, 2006.

Dated: September 20, 2006.

**Joseph T. Rannazzisi,**

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

[FR Doc. E6–16032 Filed 9–28–06; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Registration

By Notice dated May 17, 2006, and published in the **Federal Register** on May 25, 2006, (71 FR 30166), Lilly Del Caribe, Inc., Chemical Plant, Kilometer 146.7, State Road 2, Mayaguez, Puerto Rico 00680, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Dextropropoxyphene (9273), a basic class of controlled substance listed in Schedule II.

The company plans to manufacture the listed 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 Lilly Del Caribe, Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Lilly Del Caribe, 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: September 20, 2006.

**Joseph T. Rannazzisi,**

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

[FR Doc. E6–16022 Filed 9–28–06; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Manufacturer of Controlled Substances; Notice of Registration

By Notice dated June 1, 2006, and published in the **Federal Register** on June 8, 2006, (71 FR 33315), Mallinckrodt Inc., 3600 North Second Street, St. Louis, Missouri 63147, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in Schedules I and II:

| Drug | Schedule |
|---|---|
| Codeine-N-oxide (9053) | I |
| Difenoxin (9168) | I |
| Dihydromorphine (9145) | I |
| Morphine-N-oxide (9307) | I |
| Norlevorphanol (9634) | I |
| Normorphine (9313) | I |
| Tetrahydrocannabinols (7370) | I |
| Alfentanil (9737) | II |
| Amphetamine (1100) | II |
| Ecgonine (9180) | II |
| Codeine (9050) | II |
| Dextropropoxyphene, bulk (9273) | II |
| Dihydrocodeine (9120) | II |
| Diphenoxylate (9170) | II |
| Diprenorphine (9058) | II |
| Etorphine HCL (9059) | II |
| Fentanyl (9801) | II |
| Hydrocodone (9193) | II |
| Hydromorphone (9150) | II |
| Levo-alphacetylmethadol (9648) | II |
| Levorphanol (9220) | II |
| Meperidine (9230) | II |
| Methadone (9250) | II |
| Methadone intermediate (9254) | II |
| Methamphetamine (1105) | II |
| Methylphenidate (1724) | II |
| Metopon (9260) | II |
| Morphine (9300) | II |
| Nabilone (7379) | II |
| Noroxymorphone (9668) | II |
| Opium extracts (9610) | II |
| Opium fluid extract (9620) | II |
| Opium tincture (9630) | II |
| Opium, granulated (9640) | II |
| Opium, powdered (9639) | II |
| Oxycodone (9143) | II |
| Oxymorphone (9652) | II |
| Phenazocine (9715) | II |
| Remifentanil (9739) | II |
| Sufentanil (9740) | II |
| Thebaine (9333) | II |

The firm plans to manufacture the listed controlled substances for internal use and for sale to other companies.

Since the publication of the Notice of Application, it has been determined that drug code 7360 (Marihuana) is not needed as a bulk manufacturing drug code for the company. The company has subsequently withdrawn their request to add this code to their current application for registration.