

August 15, 2006, (71 FR 46921), Dade Behring Inc., 100 GBE Drive, MS514, Post Office Box 6101, Attention: RA/QS, Newark, Delaware 19714-6101, 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
Tetrahydrocannabinols (7370)	I
Benzoyllecgonine (9180)	II
Morphine (9300)	II

The company plans to produce the listed controlled substances in bulk to be used in the manufacture of reagents and drug calibrator/controls for DEA exempt products.

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 Dade Behring Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Dade Behring 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 14, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6-21870 Filed 12-21-06; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated August 7, 2006, and published in the **Federal Register** on August 15, 2006, (71 FR 46921), Dade Behring Inc., Regulatory Affairs, Quality Systems, 20400 Mariani Avenue, Cupertino, California 95014, 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
Tetrahydrocannabinols (7370)	I
Benzoyllecgonine (9180)	II
Morphine (9300)	II

The company plans to produce the listed controlled substances in bulk to be used in the manufacture of reagents and drug calibrator/controls for DEA exempt products.

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 Dade Behring Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Dade Behring 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 14, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6-21871 Filed 12-21-06; 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 November 15, 2006, 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 Remifentanyl (9739), a basic class of controlled substance listed in schedule II.

The company plans on producing this item for sale to its customers, who are final dosage manufacturers.

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 February 20, 2007.

Dated: December 14, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6-21884 Filed 12-21-06; 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 October 31, 2006, 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 14-Hydroxymorphinone (9654), a basic class of controlled substance listed in schedule II.

The company plans to manufacture the listed controlled substances in bulk as an intermediate for use in the production of another controlled substance.

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 February 20, 2007.

Dated: December 14, 2006.

Joseph T. Rannazzisi,
*Deputy Assistant Administrator, Office of
 Diversion Control, Drug Enforcement
 Administration.*
 [FR Doc. E6-21896 Filed 12-21-06; 8:45 am]
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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

**Importer of Controlled Substances
 Notice of Registration**

By Notice dated July 19, 2006, and published in the **Federal Register** on July 26, 2006, (71 FR 42417), Meridian Medical Technologies, 255 Hermelin Drive, St. Louis, Missouri 63144, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of Morphine (9300), a basic class of controlled substance listed in schedule II.

The company plans to import products for research experimentation or clinical use and analytical testing.

One objection was received; however, it has subsequently been withdrawn. DEA has considered the factors in 21 U.S.C. § 823(a) and § 952(a) and determined that the registration of Meridian Medical Technologies to import the basic class of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971, at this time. DEA has investigated Meridian Medical 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. § 952(a) and § 958(a), and in accordance with 21 CFR § 1301.34, the above named company is granted registration as an importer of the basic class of controlled substances listed.

Dated: December 14, 2006.

Joseph T. Rannazzisi,
*Deputy Assistant Administrator, Office of
 Diversion Control, Drug Enforcement
 Administration.*
 [FR Doc. E6-21895 Filed 12-21-06; 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 November 07, 2006, Organichem Corporation, 33 Riverside Avenue, Rensselaer, New York 12144, made application by renewal to the Drug Enforcement Administration (DEA) as a bulk manufacturer of the basic classes of controlled substances listed in schedule I and II:

Drug	Schedule
Marihuana (7360)	I
Tetrahydrocannabinols (7370)	I
Amphetamine (1100)	II
Methylphenidate (1724)	II
Pentobarbital (2270)	II
Hydrocodone (9193)	II
Meperidine(9230)	II
Dextropropoxyphene (9273)	II
Fentanyl (9801)	II

The company plans to manufacture bulk controlled substances for use in product development and for distribution to its customers. In reference to drug code 7360 (Marihuana), the company plans to bulk manufacture cannabindiol 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.

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 February 20, 2007.

Dated: December 14, 2006.

Joseph T. Rannazzisi,
*Deputy Assistant Administrator, Office of
 Diversion Control, Drug Enforcement
 Administration.*
 [FR Doc. E6-21886 Filed 12-21-06; 8:45 am]
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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

**Manufacturer of Controlled
 Substances Notice of Registration**

By Notice dated July 25, 2006, and published in the **Federal Register** on July 31, 2006, (71 FR 43211), 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 basic classes of controlled substances listed in schedule II:

Drug	Schedule
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Ecgonine (9180)	II
Hydrocodone (9193)	II
Morphine (9300)	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 for further manufacture or to manufacture pharmaceutical dosage forms.

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, 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.