

manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Boehringer Ingelheim Chemical 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: July 19, 2005.

**William J. Walker,**

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

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Importer of Controlled Substances; Notice of Registration**

By Notice dated April 11, 2005 and published in the **Federal Register** on April 20, 2005 (70 FR 20600), Clinical Trial Services (US), Inc., 2661 Audubon Road, Audubon, Pennsylvania 19403, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Fentanyl (9801), a basic class of controlled substance listed in Schedule II.

The company plans to import small quantities of the listed controlled substance in dosage form to conduct clinical trials.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and 952(a) and determined that the registration of Clinical Trial Services (US), Inc. to import the basic class of controlled substance 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 Clinical Trial Services (US), 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. 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 substance listed.

Dated: July 19, 2005.

**William J. Walker,**

*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 February 23, 2005, and published in the **Federal Register** on March 4, 2005, (70 FR 10679), JFC Technologies, LLC, 100 West Main Street, Bound Brook, New Jersey 08805, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic class of controlled substances listed in Schedule II:

Drug	Schedule
Diphenoxylate (9170) .....	II
Hydrocodone (9193) .....	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 JFC Technologies, LLC to manufacture the listed basic class of controlled substances is consistent with the public interest at this time. DEA has investigated JFC Technologies, 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, 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 substances listed.

Dated: July 19, 2005.

**William J. Walker,**

*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 February 23, 2005, and published in the **Federal Register** on March 4, 2005 (70 FR 10680), Lin Zhi International Inc., 687 North Pastoria Avenue, Sunnyvale, California 94085 made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic class of controlled substances listed in Schedule II:

Drug	Schedule
Oxycodone (9143) .....	II
Hydrocodone (9193) .....	II

The company plans to manufacture the listed controlled substances in bulk for use in analysis and drug test standards.

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 Lin Zhi International Inc. to manufacture the listed basic class of controlled substances is consistent with the public interest at this time. DEA has investigated Lin Zhi International 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 substances listed.

Dated: July 19, 2005.

**William J. Walker,**

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

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