

The company plans to import small quantities of the listed controlled substances for the National Institute on Drug Abuse (NIDA) for research activities.

Comments and requests for hearings on applications to import narcotic raw material are not appropriate, 72 FR 3417 (2007). Regarding all other basic classes of controlled substances, 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 Research Triangle Institute to import the basic classes 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. 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. 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 classes of controlled substances listed.

Dated: December 5, 2011.

Joseph T. Rannazzisi,

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

[FR Doc. 2011-31767 Filed 12-9-11; 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), Title 21 of the Code of Federal Regulations (CFR), this is notice that on September 15, 2011, Johnson Matthey Pharma Services, 70 Flagship Drive, North Andover, Massachusetts 01845, 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
Methylphenidate (1724)	II
Hydrocodone (9193)	II

The company plans to utilize this facility to manufacture small quantities of the listed controlled substances in bulk and to conduct analytical testing in support of the company's primary manufacturing facility in West Deptford, New Jersey. The controlled substances manufactured in bulk at this facility will be distributed to the company's customers.

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 February 10, 2012.

Dated: December 5, 2011.

Joseph T. Rannazzisi,

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

[FR Doc. 2011-31771 Filed 12-9-11; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated August 10, 2011, and published in the **Federal Register** on August 18, 2011, 76 FR 51400, 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.

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 Cambridge Isotope Lab to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Cambridge Isotope Lab 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: December 5, 2011.

Joseph T. Rannazzisi,

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

[FR Doc. 2011-31774 Filed 12-9-11; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances Notice of Registration

By Notice dated August 10, 2011, and published in the **Federal Register** on August 18, 2011, 76 FR 51401, Chemica, 316 West 130th Street, Los Angeles, California 90061, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Methamphetamine (1105), a basic class of controlled substance listed in schedule II.

The above listed controlled substance is an intermediate in the manufacture of Benzphetamine, a schedule III non-narcotic controlled substance. The company plans to utilize a bulk active pharmaceutical ingredient (API) as an intermediate for the development of another controlled substance, and further distribution to its customers. The methamphetamine will not be sold as a commercial product.

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 Chemica to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Chemica 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: December 5, 2011.

Joseph T. Rannazzisi,

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

[FR Doc. 2011-31773 Filed 12-9-11; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated August 8, 2011, and published in the **Federal Register** on August 18, 2011, 76 FR 51402, Lin Zhi International Inc., 670 Almanor Avenue, Sunnyvale, California 94085, 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
Tetrahydrocannabinols (7370)	I
3,4-Methylenedioxymethamphetamine (7405).	I
Cocaine (9041)	II
Oxycodone (9143)	II
Hydrocodone (9193)	II
Methadone (9250)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II
Morphine (9300)	II

The company plans to manufacture the listed controlled substances as bulk reagents for use in drug abuse testing.

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 classes 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 classes of controlled substances listed.

Dated: December 5, 2011.

Joseph T. Rannazzisi,

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

[FR Doc. 2011-31768 Filed 12-9-11; 8:45 am]

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DEPARTMENT OF JUSTICE

Office of Justice Programs

[OMB Number 1121-0142]

Agency Information Collection Activities; Proposed Collection; Comments Requested: Extension of a Currently Approved Collection; Victim of Crime Act, Crime Victim Assistance Grant Program, Subgrant Award Report

ACTION: 60-Day Notice of Information
Collection Under Review.

Department of Justice (DOJ), Office of Justice Programs (OJP), Office for Victims of Crime (OVC) will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The proposed information collection is published to obtain comments from the public and affected agencies. Comments are encouraged and will be accepted for "sixty days" until February 10, 2012. This process is conducted in accordance with 5 CFR 1320.10.

If you have comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact DeLano Foster (202) 616-3612, Office for Victims of Crime, Office of Justice Programs, U.S. Department of Justice, 810 7th Street NW., Washington, DC 20531.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

- Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
- Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

- Enhance the quality, utility, and clarity of the information to be collected; and
- Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

Overview of This Information Collection

(1) *Type of Information Collection:* Extension of a currently approved collection.

(2) *Title of the Form/Collection:* Victims of Crime Act, Victim Assistance Grant Program, Subgrant Award Report.

(3) *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection:* Form number: 1121-0142. Office for Victims of Crime, Office of Justice Programs, U.S. Department of Justice.

(4) *Affected public who will be asked or required to respond, as well as a brief abstract:* Primary: State government. Other: None. The VOCA, Crime Victim Assistance Grant Program, Subgrant Award Report is a required submission by state grantees, within 90 days of their awarding a subgrant for the provision of crime victim services. VOCA and the Program Guidelines require each state victim assistance office to report to OVC on the impact of the Federal funds, to certify compliance with the eligibility requirements of VOCA, and to provide a summary of proposed activities. This information will be aggregated and serve as supporting documentation for the Director's biennial report to the President and to the Congress on the effectiveness of the activities supported by these grants.

This request is for an extension of a currently approved reporting instrument, with no revisions.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond/reply:* The number of VOCA-funded victim assistance programs varies widely from State to State. A review of information currently available to this Office on the number of active victim assistance programs in 15 states selected for variance in size and population revealed that a State would be responsible for entering subgrant data for as many as 499 programs (California) to as few as 9 programs (District of Columbia).

The estimated time to enter a record via the Grants Management System is