

bulk manufacturer of a controlled substance in Schedule I or II and prior to issuing a regulation under section 1002(a) 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 section 1301.34 to Title 21, Code of Federal Regulations (CFR), notice is hereby given that on May 23, 2002, Abbott Laboratories, 1776 North Centennial Drive, McPherson, Kansas 67460-1247, made application by renewal to the Drug Enforcement Administration to be registered as an importer of remifentanil (9739), a basic class of controlled substance listed in Schedule II.

The firm plans to import the remifentanil to manufacture Ultiva for the U.S. market.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substance may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.43 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections or requests for a hearing may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than September 18, 2002.

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 at 40 F.R. 43745-46 (September 23, 1975), all applicants for registration to import a basic class of any controlled substance 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 U.S.C. 958(a), 21 U.S.C. 823(a), and 21 CFR 1301.34(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: July 29, 2002.

Laura M. Nagel,
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 March 7, 2002, and published in the **Federal Register** on March 18, 2002, (67 FR 12049), Chemic Laboratories, Inc., 480 Neponset Street, Building 7C, Canton, Massachusetts 02021, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of cocaine (9041), a basic class of controlled substance listed in Schedule II.

The firm plans to bulk manufacture small quantities of cocaine derivative for a customer.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, section 823(a) and determined that the registration of Chemic Laboratories, Inc. to manufacture is consistent with the public interest at this time. DEA has investigated Chemic Laboratories, Inc. to ensure that the company's registration is consistent with the public interest. These investigations have 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 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic class of controlled substance listed above is granted.

Dated: July 29, 2002.

Laura M. Nagel,
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 19, 2002, and published in the **Federal Register** on March 12, 2002, (67 FR 11141), Chiragene, Inc., Technology Center of New Jersey, 661 Highway One, North Brunswick, New Jersey 08902, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as

be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
N-Ethylamphetamine (1475)	I
2,5-Dimethoxyamphetamine (7396)	I
3,4-Methylenedioxymethamphetamine (7400)	I
4-Methoxyamphetamine (7411)	I
Amphetamine (1100)	II
Methylphenidate (1724)	II
Morphine (9300)	II
Fentanyl (9801)	II

The firm plans to manufacture the listed controlled substances to supply their customers.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, section 823(a) and determined that the registration of Chiragene, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Chiragene, Inc. on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, 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 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: July 29, 2002.

Laura M. Nagel,
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 March 27, 2002, and published in the **Federal Register** on April 10, 2002, (67 FR 17470), Organix Inc., 240 Salem Street, Woburn, Massachusetts 01801, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as

a bulk manufacturer of cocaine (9041), a basic class of controlled substance listed in Schedule II.

The firm plans to manufacture a derivative of cocaine in gram quantities for validation of synthetic procedures.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Organix, Inc. to manufacture is consistent with the public interest at this time. DEA has investigated Organix, Inc. to ensure that the company's registration is consistent with the public interest. This investigation 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 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic class of controlled substance listed above is granted.

Dated: July 29, 2002.

Laura M. Nagel,

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 March 29, 2001, and published in the **Federal Register** on April 6, 2001, (66 FR 18309) Ganes Chemicals Inc., which has changed its name to Siegfried (USA), Inc., 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 below:

Drug	Schedule
Amphetamine (1100)	II
Methylphenidate (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Glutethimide (2250)	II
Methadone (9250)	II
Methadone-intermediate (9254)	II

Drug	Schedule
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II

The firm plans to manufacture the listed controlled substances for distribution as bulk products to its customers.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, section 823(a) and determined that the registration of Siegfried (USA), Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Siegfried (USA), on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, 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 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: July 29, 2002.

Laura M. Nagel,

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

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DEPARTMENT OF JUSTICE

Office of Justice Programs

Agency Information Collection Activities: Proposed Collection; Comment Request

ACTION: 30-day notice of information collection under review: reinstatement, with change, of a previously approved collection for which approval has expired, survey of inmates in State and Federal correctional facilities, 2003.

The Department of Justice (DOJ), Office of Justice Programs, has submitted 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. This proposed information collection was previously published in the **Federal Register** Volume 67, Number 106, page 38295 on June 3, 2002, allowing for a 60-day public comment period.

The purpose of this notice is to allow an additional 30 days for public comment until September 18, 2002. This process is conducted in accordance with 5 CFR 1320.10.

Written comments and/or suggestions regarding the items contained in this notice, especially the estimated public burden and associated response time, should be directed to the Office of Management and Budget, Office of Information and Regulatory Affairs, Attention Department of Justice Desk Officer, Washington, DC 20503. Additionally, comments may be submitted to OMB via facsimile to (202) 395-7285.

Request 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:

(1) Evaluate whether the collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(3) Enhance the quality, utility and clarity of the information to be collected; and

(4) 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:* Reinstate, with Change, of a Previously Approved Collection for which Approval has Expired.

(2) *Title of the Form/Collection:* Survey of Inmates in State and Federal Correctional Facilities, 2003.

(3) *Agency Form Number and the applicable component of the Department sponsoring the collection:* Form: NPS-25 CAPI Instrument; NPS-13 Sampling Questionnaire; and NPS-27, Inmate letter. Corrections Unit, Bureau of Justice Statistics, Office of