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: November 20, 2000.

John H. King,

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

[FR Doc. 00–30935 Filed 12–4–00; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By notice dated August 14, 2000 and published in the **Federal Register** on August 23, 2000, (65 FR 51330), Applied Science Labs, Inc., A Division of Alltech Associates, Inc., 2701 Carolean Industrial Drive, P.O. Box 440, State College, Pennsylvania 16801, made application by renewal to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Heroine (9200)	I
Cocaine (9041)	II
Codeine (9050)	II
Meperidine (9230)	II
Methadone (9250)	II
Morphine (9300)	II

The firm plans to import these controlled substances for the manufacture of reference standards.

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 Applied Science Labs, Inc., to import the listed 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 Applied Science Labs, 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 section 1008(a) of the Controlled Substances Import and Export Act and in accordance with title 21, Code of Federal Regulations, § 1301.34, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

November 20, 2000.

John H. King,

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

[FR Doc. 00–30938 Filed 12–4–00; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importation of Controlled Substances; Notice of Registration

By notice dated August 8, 2000, and published in the **Federal Register** on August 23, 2000 (65 FR 51331), Calbiochem-Novabiochem Corporation, 10394 Pacific Center Court, Attn: Receiving Inspector, San Diego, California 92121–4340, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Tetrahydrocannabinols (7370)	I
Mescaline (7381)	I
Phencyclidine (7471)	II
Phenylaceton (8501)	II
Cocaine (9041)	II

The firm plans to import small quantities of the listed controlled substances to make reagents for distribution to the biomedical research community.

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 Calbiochem-Novabiochem Corporation 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 Calbiochem-Novabiochem Corporation 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 section 1008(a) of the Controlled Substance Import and Export Act and in accordance with the Title 21, Code of Federal Regulations, § 1301.34, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: November 20, 2000.

John H. King,

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

[FR Doc. 00–30939 Filed 12–4–00; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated June 29, 2000, and published in the **Federal Register** on July 14, 2000, (65 FR 43785), 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 the basic classes of controlled substances listed below:

Drug	Schedule
Methaqualone (2565)	I
Dimethyltryptamine (7435)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Phencyclidine (7471)	II
Phenylacetone (8501)	II
Cocaine (9041)	II
Codeine (9050)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Benzoylecgonine (9180)	II
Methadone (9250)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II
Morphine (9300)	II
Fentanyl (9801)	II

The firm plans to manufacture small quantities of the listed controlled substances to produce isotope labeled standards for drug analysis.

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 Cambridge Isotope Lab to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Cambridge Isotope Lab on a regular basis to ensure that its 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: November 20, 2000.

John H. King,

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

[FR Doc. 00–30932 Filed 12–4–00; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated June 14, 2000, and published in the **Federal Register** on June 26, 2000, (65 FR 39430), Celgene Corporation, 7 Powder Horn Drive, Warren, New Jersey 07059, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of methylphenidate (1724), a basic class of controlled substance listed in Schedule II.

The firm plans to manufacture methylphenidate for product research and development.

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 Celgene Corporation to manufacture methylphenidate is consistent with the public interest at this time. DEA has investigated the Celgene Corporation 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 class of controlled substance listed above is granted.

Dated: October 10, 2000.

John H. King,

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

[FR Doc. 00–30933 Filed 12–4–00; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By notice dated June 29, 2000, and published in the **Federal Register** on July 14, 2000, (65 FR 43785), Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Building 18, Chattanooga, Tennessee 37409, 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
Methamphetamine (1105)	II

No comments or objections have been received. The firm plans to bulk manufacture amphetamine and methamphetamine to produce products for distribution to its customers.

DEA has considered the factors in title 21, United States Code, section 823(a) and determined that the registration of Chattem Chemicals, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Chattem Chemicals, 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: November 20, 2000.

John H. King,

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

[FR Doc. 00–30937 Filed 12–4–00; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

The Church of the Living Tree; Notice of Withdrawal of Denial of Application

On November 4, 1999, the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration (DEA) issued an Order to Show Cause proposing to deny The Church of the Living Tree's application for registration as a manufacturer of the controlled substance marijuana. Effective August 18, 2000, the DEA Administrator, in his final order published in the **Federal Register**, 65 FR 50567, denied the application for registration, noting that The Church of the Living Tree had not responded to the Order to Show Cause.

By letter dated August 11, 2000, a representative of The Church of the Living Tree advised that he had in fact properly submitted a timely request for hearing. Photocopies of a United States Postal Service Receipt for Certified Mail and Domestic Return Receipt indicating delivery accompanied the letter.

Inasmuch as it appears that The Church of the Living Tree timely requested a hearing in this matter, the final order should not have issued. The Deputy Administrator of the Drug Enforcement Administration therefore orders that the final order published August 18, 2000, at 65 Fed. Reg. 50567 be, and it hereby is, rescinded, and this matter is hereby remanded to the Office of Administrative Law Judges, Drug Enforcement Administration, for further appropriate proceedings. This order is effective December 5, 2000.