

interest at this time. DEA has investigated S & B Pharma 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(a), 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: April 17, 2013.

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

[FR Doc. 2013-09532 Filed 4-22-13; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration; Alltech Associates, Inc.

By Notice dated November 14, 2012 and published in the **Federal Register** on November 23, 2012, 77 FR 70188, Alltech Associates, Inc., 2051 Waukegan Road, Deerfield, Illinois 60015, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
2C-T-2 (2-(4-Ethylthio-2,5-dimethoxyphenyl)ethanamine) (7385)	I
2C-1 (2-(4-Iodo-2,5-dimethoxyphenyl)ethanamine) (7518)	I
2C-C (2-(4-Chloro-2,5-dimethoxyphenyl)ethanamine) (7519)	I

The company plans to manufacture high purity drug standards used for analytical applications only in clinical, toxicological, and forensic laboratories.

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 Alltech Associates, Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Alltech Associates, 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(a), 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: April 16, 2013.

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

[FR Doc. 2013-09531 Filed 4-22-13; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration; PCAS-Nanosyn, LLC

By Notice dated January 15, 2013, and published in the **Federal Register** on January 30, 2013, 78 FR 6350, PCAS-Nanosyn, LLC, 3331-B Industrial Drive, Santa Rosa, California 95403, 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
Methamphetamine (1105)	II
Methylphenidate (1724)	II
Phencyclidine (7471)	II
Codeine (9050)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Hydrocodone (9193)	II
Methadone (9250)	II
Morphine (9300)	II
Oxymorphone (9652)	II
Fentanyl (9801)	II

The company is a contract manufacturer. At the request of the company's customers, it manufactures derivatives of controlled substances in bulk form only.

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 PCAS-Nanosyn, LLC., to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has

investigated PCAS-Nanosyn, 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(a), 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: April 16, 2013.

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

[FR Doc. 2013-09529 Filed 4-22-13; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration; Cayman Chemical Company

By Notice dated November 14, 2012, and published in the **Federal Register** on November 23, 2012, 77 FR 70188, Cayman Chemical Company, 1180 East Ellsworth Road, Ann Arbor, Michigan 48108, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances: