DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Manufacturer of Controlled Substances Registration: Boehringer Ingelheim Chemical, Inc.

ACTION: Notice of registration.

SUMMARY: Boehringer Ingelheim Chemical, Inc. applied to be registered as a manufacturer of certain basic classes of controlled substances. The DEA grants Boehringer Ingelheim Chemical, Inc. registration as a manufacturer of the controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated May 28, 2014, and published in the **Federal Register** on June 4, 2014, 79 FR 32321, Boehringer Ingelheim Chemical, Inc., 2820 N. Normandy Drive, Petersburg, Virginia 23805–9372, applied to be registered as a manufacturer of certain basic classes of controlled substances. No comments or objections were submitted to this notice.

The Drug Enforcement Administration (DEA) has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Boehringer Ingelheim Chemical, Inc. to manufacture 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. The DEA investigated the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing 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:

Controlled substance	Schedule
Amphetamine (1100)	II
Methylphenidate (1724)	II
Methadone (9250)	II
Methadone Intermediate (9254)	II

The company plans to manufacture the listed controlled substances in bulk for sale to its customers for formulation into finished pharmaceuticals. In reference to methadone intermediate (9254) the company plans to produce methadone HCL active pharmaceutical ingredients (APIs) for sale to its customers.

Dated: January 9, 2015.

Joseph T. Rannazzisi,

Deputy Assistant Administrator.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Manufacturer of Controlled Substances Registration: S & B Pharma, Inc.

ACTION: Notice of registration.

SUMMARY: S & B Pharma, Inc. applied to be registered as a manufacturer of certain basic classes of controlled substances. The DEA grants S & B Pharma, Inc. registration as a manufacturer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated July 1, 2014, and published in the **Federal Register** on July 8, 2014, 79 FR 38564, S & B Pharma, Inc., DBA Norac Pharma, 405 South Motor Avenue, Azusa, California 91702–3232 applied to be registered as a manufacturer of certain basic classes of controlled substances. No comments or objections were submitted to this notice.

The Drug Enforcement Administration (DEA) has considered the factors in 21 U.S.C. 823(a) and determined that the registration of S & B Pharma, Inc. to manufacture 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. The DEA investigated the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing 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:

Controlled Substance	Schedule
Gamma Hydroxybutyric Acid (2010).	I
Tetrahydrocannabinols (7370)	I
Methamphetamine (1105)	II
Pentobarbital (2270)	II
Nabilone (7379)	II
4-Anilino-N-phenethyl-4-piperidine (ANPP) (8333).	II
Tapentadol (9780)	II
Fentanyl (9801)	II

The company plans to manufacture the listed controlled substances in bulk for use in product development and for commercial sales to its customers.

Dated: January 9, 2015.

Joseph T. Rannazzisi,

Deputy Assistant Administrator.

[FR Doc. 2015–01287 Filed 1–23–15; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Manufacturer of Controlled Substances Registration: Research Triangle Institute

ACTION: Notice of registration.

SUMMARY: Research Triangle Institute applied to be registered as a manufacturer of certain basic classes of controlled substances. The DEA grants Research Triangle Institute registration as a manufacturer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated July 2, 2014, and published in the **Federal Register** on July 14, 2014, 79 FR 40781, Research Triangle Institute, Kenneth S. Rehder, Ph.D., Hermann Building East Institute Drive, P.O. Box 12194, Research Triangle Park, North Carolina 27709, applied to be registered as a manufacturer of certain basic classes of controlled substances. No comments or objections were submitted to this notice.

The Drug Enforcement Administration (DEA) has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Research Triangle Institute to manufacture 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. The DEA investigated the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing the company's background and history.