

is notice that on October 4, 2019, S & B Pharma, Inc., DBA Norac Pharma, 405

South Motor Avenue, Azusa, California 91702-3232 applied to be registered as

a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid .....	7360	I
Tetrahydrocannabinols .....	7370	I
Amphetamine .....	1100	II
Methamphetamine .....	1105	II
Lisdexamfetamine .....	1205	II
Methylphenidate .....	1724	II
Pentobarbital .....	2270	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. In reference to drug code 7360 (marihuana) and 7370 (tetrahydrocannabinols), the company plans to bulk manufacture both as synthetic substances. No other activity for these drug codes is authorized for this registration.

Dated: November 5, 2019.

**William T. McDermott,**  
Assistant Administrator.

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-549]

**Importer of Controlled Substances Application: Mylan Technologies Inc.**

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before December 23, 2019. Such persons may also file a written request for a hearing on the application on or before December 23, 2019.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701

Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on October 16, 2019, Mylan Technologies Inc., 110 Lake Street, Saint Albans, Vermont 054780 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Methylphenidate .....	1724	II
Fentanyl .....	9801	II

The company plans to import the listed controlled substances in finished dosage form (FDF) from foreign sources for analytical testing and clinical trials in which the foreign FDF will be compared to the company's own domestically-manufactured FDF. This analysis is required to allow the company to export domestically manufactured FDF to foreign markets.

Dated: November 8, 2019.

**William T. McDermott,**  
Assistant Administrator.

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-540]

**Bulk Manufacturer of Controlled Substances Application: Chattem Chemicals**

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before January 21, 2020.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.33(a), this is notice that on May 17, 2019, Chattem Chemicals, 3801 Saint Elmo Avenue, Chattanooga, Tennessee 37409 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid .....	2010	I
Marihuana .....	7360	I
Tetrahydrocannabinols .....	7370	I