Billing Code 4110-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-545]

Bulk Manufacturer of Controlled Substances Application: S&B Pharma, 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 on or before [INSERT 60 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER].

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

SUPPLEMENTARY INFORMATION:

In accordance with 21 CFR 1301.33(a), this 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
Telrahydrocannabinois	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 (tetrahydrocannabinois), the company plans to bulk manufacture both as synthetic substances. No other activity for these dug codes is authorized for this registration.

Dated: November 5, 2019.

William T. McDermott,
Assistant Administrator.

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