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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-529]

Bulk Manufacturer of Controlled Substances Application: Patheon API Manufacturing, 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 [INSERT 60 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER].

ADDRESS: 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 March 15, 2019, Patheon API Manufacturing, Inc., 309 Delaware Street, Greenville, South Carolina 29605 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled Substance	Drug Code	Schedule
Thebaine	9333	II
Noroxymorphone	9668	II
Gamma Hydroxybutyric Acid	2010	I
Alpha-methyltryptamine	7432	I

The company plans to bulk manufacture the listed controlled substances as an Active Pharmaceutical Ingredient for supply to its customers.

Dated: November 5, 2019.

William T. McDermott,
Assistant Administrator.

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