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.

[FR Doc. 2019–25402 Filed 11–21–19; 8:45 am]

BILLING CODE 4410–09–P

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

[FR Doc. 2019–25405 Filed 11–21–19; 8:45 am]

BILLING CODE 4410–09–P

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 Morrissette 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

Controlled substance	Drug code	Schedule
4-Methoxyamphetamine	7411	I
Dihydromorphine	9145	I
Amphetamine	1100	II
Methamphetamine	1105	II
Lisdexamfetamine	1205	II
Methylphenidate	1724	II
Cocaine	9041	II
Codeine	9050	II
Dihydrocodeine	9120	II
Oxycodone	9143	II
Hydromorphone	9150	II
Ecgonine	9180	II
Hydrocodone	9193	II
Levorphanol	9220	II
Methadone	9250	II
Methadone intermediate	9254	II
Morphine	9300	II
Oripavine	9330	II
Thebaine	9333	II
Oxymorphone	9652	II
Noroxymorphone	9668	II
Tapentadol	9780	II
Fentanyl	9801	II

The company plans to manufacturer the listed controlled substances in bulk for distribution and sale to its customers.

In reference to drug codes 7360 (marihuana) and 7370 (tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetics. No other activities for these drug codes are authorized for this registration.

Dated: November 5, 2019.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2019–25400 Filed 11–21–19; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

[OMB Number 1117–0053]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Extension of a Previously Approved Collection: Leadership Engagement Survey

AGENCY: Department of Justice, Drug Enforcement Administration.

ACTION: 30-Day notice.

SUMMARY: The Department of Justice (DOJ), Drug Enforcement Administration (DEA), will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The proposed information collection was previously published in the **Federal Register** allowing for a 60 day comment period.

DATES: Comments are encouraged and will be accepted for 30 days until December 23, 2019.

FOR FURTHER INFORMATION CONTACT: If you have additional comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact Chiwoniso S. Gurira (Choni), Senior Personnel Psychologist, Drug Enforcement Administration, 8701 Morrissette Drive, Springfield, VA 22152. Written comments and/or suggestions may also be directed to the Office of Management and Budget, Office of Information and Regulatory Affairs, Attention Department of Justice Desk Officer, Washington, DC 20530 or send to *OIRA_submission@omb.eop.gov.*

SUPPLEMENTARY INFORMATION: This process is conducted in accordance with 5 CFR 1320.10. Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

—Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the Bureau of Justice Statistics, including whether the information will have practical utility;

—Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

—Evaluate whether and if so how the quality, utility, and clarity of the information to be collected can be enhanced; and

—Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses.

Overview of This Information Collection

1. *Type of Information Collection:* Extension of a previously approved collection for which approval has expired.

2. *The Title of the Form/Collection:* Leadership Engagement Survey.

3. *The agency form number, if any and the applicable component of the Department sponsoring the collection:* Online survey.

4. *Affected public who will be asked or required to respond, as well as a brief abstract:*

Primary: Drug Enforcement Administration contractors and Task Force Officers.

Other: None.

5. *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* It is estimated that 5000 respondents will complete the survey in approximately 20 minutes.

6. *An estimate of the total public burden (in hours) associated with the collection:* The estimated public burden associated with this collection is 1667 hours. It is estimated that applicants