Issued: November 20, 2019.

William Bishop,

Supervisory Hearings and Information Officer.

[FR Doc. 2019–25500 Filed 11–20–19; 11:15 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–550]

Importer of Controlled Substances Application: Janssen Pharmaceuticals, 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 9, 2019, Janssen Pharmaceuticals, Inc., 1440 Olympic Drive, Athens, Georgia 30601–1645 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Thebaine	9333	II
Concentrate of Poppy Straw	9670	II
Tapentadol	9780	II

The company plans to import intermediate forms of tapentadol (9780) and thebaine (9333) for further manufacturing prior to distribution to its customers. The company plans to import concentrate of poppy straw (9670) to bulk manufacture other controlled substances. No other activity for these drug codes is authorized for this registration.

Dated: November 8, 2019.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2019–25407 Filed 11–21–19; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–551]

Importer of Controlled Substances Application: Epic Pharma, LLC

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 September 13, 2019, Epic Pharma, LLC, 227–15 North Conduit Avenue, Laurelton, New York 11413, applied to be registered as an importer of the following basic class of controlled substance:

Controlled substance	Drug code	Schedule
Methadone	9250	II

The company plans to import the listed controlled substance for research and analytical purposes.

Dated: November 8, 2019.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2019–25406 Filed 11–21–19; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–528]

Importer of Controlled Substances Application: Fresenius Kabi USA, LLC

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 August 7, 2019 Fresenius Kabi USA, LLC, 3159 Staley Road, Grand Island, New York 14072–

2028 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Remifentanil	9739	II

The company plans to import the listed controlled substance for bulk manufacture.

Dated: October 23, 2019.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2019–25403 Filed 11–21–19; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–538]

Importer of Controlled Substances Application: GE Healthcare

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 July 15, 2019, GE Healthcare, 3350 North Ridge Avenue, Arlington Heights, Illinois 60004–1412 applied to be registered as an importer of the following basic class of controlled substance:

Controlled substance	Drug code	Schedule
Cocaine	9041	II

The company plans to import small quantities of Ioflupane, in the form of three separate analogues of cocaine, to validate production and quality control systems, for a reference standard, and for producing material for a future investigational new drug (IND) submission. Supplies of this particular controlled substance are not available in the form needed within the current domestic supply of the United States.

Dated: November 7, 2019.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2019–25404 Filed 11–21–19; 8:45 am]

BILLING CODE 4410–09–P

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 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 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.

[FR Doc. 2019–25401 Filed 11–21–19; 8:45 am]

BILLING CODE 4410–09–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 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