

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 June 24, 2020, Cedarburg Pharmaceuticals, 870 Badger Circle, Grafton, Wisconsin 53024-0000, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substances:

Controlled substance	Drug code	Schedule
Tetrahydrocannabinols ..	7370	I
Methylphenidate	1724	II
Nabilone	7379	II
4-Anilino-N-phenethyl-4-piperidine (ANPP).	8333	II
Fentanyl	9801	II

The company plans to manufacture bulk active pharmaceutical ingredients (API) for distribution to its customers. In reference to drug code 7370 (Tetrahydrocannabinols) the company plans to bulk manufacture as synthetic. No other activity for this drug code is authorized for this registration.

William T. McDermott,
Assistant Administrator.

[FR Doc. 2020-17434 Filed 8-10-20; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-695]

Importer of Controlled Substances Application: Epic Pharma, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Epic Pharma, LLC applied to be registered as an importer of the following basic class(es) of a controlled substance: Methadone.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before September 10, 2020. Such persons may also file a written request for a hearing on the application on or before September 10, 2020.

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 request 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 July 10, 2020, Epic Pharma, LLC, 227-15 North Conduit Avenue, Laurelton, New York 11413 applied to be registered as an importer of the following basic class(es) of a controlled substance:

Controlled substance	Drug code	Schedule
Methadone	9250	II

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

William T. McDermott,
Assistant Administrator.

[FR Doc. 2020-17432 Filed 8-10-20; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-701]

Bulk Manufacturer of Controlled Substances Application: Purisys, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Purisys, LLC applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substances: Lysergic acid diethylamide and Pentobarbital.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before October 13, 2020.

ADDRESS: 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 July 15, 2020, Purisys,

LLC, 1550 Olympic Drive, Athens, Georgia 30601-1602, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substances:

Controlled substance	Drug code	Schedule
Lysergic acid diethylamide.	7315	I
Pentobarbital	2270	II

The company plans to manufacture the above-listed controlled substances as analytical reference standards and clinical trial material for distribution to its customers. No other activities for these drug codes are authorized for this registration.

William T. McDermott,
Assistant Administrator.

[FR Doc. 2020-17438 Filed 8-10-20; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-697]

Importer of Controlled Substances Application: GE Healthcare

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: GE Healthcare applied to be registered as an importer of the following basic class(es) of a controlled substance: Cocaine.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before September 10, 2020. Such persons may also file a written request for a hearing on the application on or before September 10, 2020.

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.