

Controlled substance	Drug code	Schedule
Metazocine	9240	II
Methadone	9250	II
Methadone intermediate	9254	II
Metopon	9260	II
Dextropropoxyphene, bulk (non-dosage forms)	9273	II
Morphine	9300	II
Oripavine	9330	II
Thebaine	9333	II
Dihydroetorphine	9334	II
Opium, raw	9600	II
Opium extracts	9610	II
Opium fluid extract	9620	II
Opium tincture	9630	II
Opium, powdered	9639	II
Opium, granulated	9640	II
Levo-alphaacetylmethadol	9648	II
Opium poppy	9650	II
Oxymorphone	9652	II
Poppy Straw Concentrate	9670	II
Phenazocine	9715	II
Piminodine	9730	II
Racemethorphan	9732	II
Racemorphan	9733	II
Alfentanil	9737	II
Remifentanil	9739	II
Sufentanil	9740	II
Carfentanil	9743	II
Tapentadol	9780	II
Bezitramide	9800	II
Fentanyl	9801	II
Moramide-intermediate	9802	II

The company plans to import small quantities of the listed controlled substances for the National Institute on Drug Abuse (NIDA) for research activities. The company plans to import analytical reference standards for distribution to its customers for research and analytical purposes.

Dated: July 16, 2019.

John J. Martin,
Assistant Administrator.

[FR Doc. 2019-16167 Filed 7-29-19; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Application: Cambrex Charles City

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 August 29, 2019. Such persons may also file a written request for a hearing on the application on or before August 29, 2019.

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.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on May 15, 2019, Cambrex Charles City, 1205 11th Street, Charles City, Iowa 50616 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
4-Anilino-N-phenethyl-4-piperidine (ANPP)	8333	II
Phenylacetone	8501	II
Coca Leaves	9040	II
Opium, raw	9600	II
Poppy Straw Concentrate	9670	II

The company plans to import the listed controlled substances for internal

use, and to manufacture bulk intermediates for sale to its customers.

Dated: July 16, 2019.

John J. Martin,
Assistant Administrator.

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BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Application: Catalent Pharma Solutions, 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 August 29, 2019. Such persons may also file a written request for a hearing on the application on or before August 29, 2019.

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.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on May 31, 2019, Catalent Pharma Solutions, LLC, 3031 Red Lion Road, Philadelphia, Pennsylvania 19114 applied to be registered as an importer of the following basic class of controlled substance:

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid.	2010	I

The company plans to import finished dosage unit products containing gamma-

hydroxybutyric acid for clinical trials, research, and analytical activities.

Dated: July 16, 2019.
John J. Martin,
Assistant Administrator.
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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Application: Xcelience

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 August 29, 2019. Such persons may also file a written request for a hearing on the application on or before August 29, 2019.

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 May 2, 2019, Xcelience, 4901 West Grace Street, Tampa, Florida 33607, applied to be registered as an importer of the following basic class of controlled substance:

Controlled substance	Drug code	Schedule
Amphetamine	1100	II

The company plans to import the listed controlled substance in finished dosage form for clinical trials, research and analytical purposes.

Dated: July 16, 2019.
John J. Martin,
Assistant Administrator.
 [FR Doc. 2019-16168 Filed 7-29-19; 8:45 am]
BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Application: Nostrum Laboratories, 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 August 29, 2019. Such persons may also file a written request for a hearing on the application on or before August 29, 2019.

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.

SUPPLEMENTARY INFORMATION:

In accordance with 21 CFR 1301.34(a), this is notice that on March 13, 2019, Nostrum Laboratories, Inc., 705 East Mulberry Street, Bryan, Ohio 43506 applied to be registered as an importer of the following basic class of controlled substance:

Controlled substance	Drug code	Schedule
Marihuana Extract	7350	I
Marihuana	7360	I

The company plans to import the listed controlled substances for research and new drug development. Approval of permit applications will occur only when the registrant's business activity is

consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of FDA approved or non-approved finished dosage forms for commercial sale.

Dated: July 16, 2019.
John J. Martin,
Assistant Administrator.
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