

reference to drug codes 7360 (marihuana), and 7370 (tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetic cannabidiol and tetrahydrocannabinol. No other activity for these drug codes is authorized for this registration.

William T. McDermott,
Assistant Administrator.

[FR Doc. 2020-20596 Filed 9-17-20; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-705]

Importer of Controlled Substances Application: Fisher Clinical Services, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Fisher Clinical Services, Inc. has applied to be registered as an importer of basic class(es) of controlled substances. Refer to Supplemental Information listed below for further drug information.

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 19, 2020. Such persons may also file a written request for a hearing on the application on or before October 19, 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.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on August 3, 2020, Fisher Clinical Services, Inc., 7554 Schantz Road, Allentown, Pennsylvania 18106-9032, applied to be registered as an importer of the following basic class(es) of controlled substances:

Controlled substance	Drug code	Schedule
Psilocybin	7437	I
Methylphenidate	1724	II
Levorphanol	9220	II
Noroxymorphone	9668	II
Tapentadol	9780	II

The company plans to import the listed controlled substances for clinical trials only.

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 the Food and Drug Administration approved or non-approved finished dosage forms for commercial sale.

William T. McDermott,
Assistant Administrator.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-706]

Bulk Manufacturer of Controlled Substances Application: Cambrex High Point, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Cambrex High Point, Inc., has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to Supplemental Information listed below for further drug information.

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 November 17, 2020. Such persons may also file a written request for a hearing on the application on or before November 17, 2020.

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 July 27, 2020, Cambrex High Point, Inc., 4180 Mendenhall Oaks Parkway, High Point, North Carolina 27265-8017, applied to be registered as an bulk manufacturer of the following basic class(es) of controlled substances:

Controlled substance	Drug code	Schedule
Oxymorphone	9652	II
Noroxymorphone	9668	II

The company plans to manufacture the above-listed controlled substances in bulk for distribution to its customers. No other activities for these drug codes are authorized for this registration.

William T. McDermott,
Assistant Administrator.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-721]

Bulk Manufacturer of Controlled Substances Application: Nalas Engineering Services, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Nalas Engineering Services, Inc. has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to Supplemental Information listed below for further drug information.

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 November 17, 2020. Such persons may also file a written request for a hearing on the application on or before November 17, 2020.

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 August 13, 2020, Nalas Engineering Services, Inc., 85 Westbrook Road, Centerbrook, Connecticut 06409, applied to be registered as an bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Thebaine	9333	II