

Issued: October 1, 2020.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2020–22027 Filed 10–5–20; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–727]

Bulk Manufacturer of Controlled Substances Application: Halo Pharmaceutical, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Halo Pharmaceutical, 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 December 7, 2020. Such persons may also file a written request for a hearing on the application on or before December 7, 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 September 3, 2020, Halo Pharmaceutical Inc, 30 North Jefferson Road, Whippany, New Jersey 07981, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substances	Drug codes	Schedule
Dihydromorphine	9145	I
Hydromorphone	9150	II

The company plans to manufacture Hydromorphone (9150) for distribution to its customers. Dihydromorphine (9145) is an intermediate in the manufacture of Hydromorphone and is not for commercial distribution. No other activity for these drug codes is authorized for this registration.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–22076 Filed 10–5–20; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–726]

Bulk Manufacturer of Controlled Substances Application: S&B Pharma, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: S&B Pharma, LLC 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 December 7, 2020. Such persons may also file a written request for a hearing on the application on or before December 7, 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 25, 2020, S&B Pharma, LLC, 405 South Motor Avenue, Azusa, California 91702, applied to be registered as an bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid	2010	I
Marihuana	7360	I
Tetrahydrocannabinols	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 (Tetrahydrocannabinols), the company plans to bulk manufacture both as synthetic substances. No other

activity for these drug codes is authorized for this registration.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–22075 Filed 10–5–20; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–725]

Importer of Controlled Substances Application: Wildlife Laboratories, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Wildlife Laboratories, LLC has applied to be registered as an importer 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 5, 2020. Such persons may also file a written request for a hearing on the application on or before November 5, 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 September 4, 2020, Wildlife Laboratories, LLC, 1230 W Ash Street, Unit D, Windsor, Colorado

80550–4677, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Etorphine HCL	9059	II
Thiafentanil	9729	II

The company plans to import the listed controlled substances for distribution to its customers. No other activity for these drug codes is authorized for this registration.

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.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–22069 Filed 10–5–20; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–722]

Bulk Manufacturer of Controlled Substances Application: Eli Elsohly Laboratories

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Eli Elsohly Laboratories 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 December 7, 2020. Such persons may also file a written request for a hearing on the application on or before December 7, 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 17, 2020, Eli Elsohly Laboratories, 5 Industrial Park Drive, Oxford, Mississippi 38655, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substances	Drug code	Schedule
Marihuana Extract	7350	I
Marihuana	7360	I
Tetrahydrocannabinols	7370	I
Dihydromorphone	9145	I
Amphetamine	1100	II
Methamphetamine	1105	II
Cocaine	9041	II
Codeine	9050	II
Dihydrocodeine	9120	II
Oxycodone	9143	II
Hydromorphone	9150	II
Ecgonine	9180	II
Hydrocodone	9193	II
Morphine	9300	II
Thebaine	9333	II
Oxymorphone	9652	II

The company plans to manufacture the listed controlled substances for product development and reference standards. In reference to drug codes 7360 (marihuana) and 7370 (THC), the company plans to isolate these controlled substances from procured 7350 (marihuana extract). In reference to drug code 7360, no cultivation activities are authorized for this registration. No

other activities for these drug codes are authorized for this registration.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–22068 Filed 10–5–20; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Federal Bureau of Investigation

Meeting of the Compact Council for the National Crime Prevention and Privacy Compact

AGENCY: Federal Bureau of Investigation, Department of Justice.