The company plans to import the listed control substances for clinical trials, research and development, analytical purposes, and 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 Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

Brian S. Besser,

Acting Assistant Administrator.

[FR Doc. 2021–23036 Filed 10–21–21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–912]

Bulk Manufacturer of Controlled Substances Application: Groff NA Hemplex LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Groff NA Hemplex LLC has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTARY 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 21, 2021. Such persons may also file a written request for a hearing on the application on or before December 21, 2021.

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 August 16, 2021, Groff NA Hemplex LLC, 100 Redco Avenue, Suite A, Red Lion, Pennsylvania 17356–1436, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Marihuana Extract	7350	I
Marihuana	7360	I
Tetrahydrocannabinols	7370	I

The company is federally authorized to conduct cultivation activities in order to bulk manufacture the listed controlled substances for internal use and for sale to federally registered research investigators. No other activities for these drug codes are authorized for this registration.

Brian S. Besser,

Acting Assistant Administrator.

[FR Doc. 2021–23034 Filed 10–21–21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–914]

Bulk Manufacturer of Controlled Substances Application: Organic Consultants, LLC. DBA Cascade Chemistry

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Organic Consultants, LLC. DBA Cascade Chemistry, has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to Supplementary 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 21, 2021. Such persons may also file a written request for a hearing on the application on or before December 21, 2021.

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 September 9, 2021, Organic Consultants, LLC. DBA Cascade Chemistry, 90 North Polk Street, Suite 200, Eugene, Oregon 97402–4109, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Amphetamine	1100	II
Methylphenidate	1724	II
Nabilone	7379	II
Codeine	9050	II
Oxycodone	9143	II
Hydromorphone	9150	II
Hydrocodone	9193	II
Methadone	9250	II
Methadone intermediate	9254	II
Morphine	9300	II
Thebaine	9333	II
Oxymorphone	9652	II
Fentanyl	9801	II

The company plans to bulk manufacture small quantities of the listed controlled substances for internal use or for sale as analytical reference standard materials to its customers. No other activities for these drug codes are authorized for this registration.

Brian S. Besser,

Acting Assistant Administrator.

[FR Doc. 2021–23038 Filed 10–21–21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–911]

Importer of Controlled Substances Application: Novitium Pharma LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Novitium Pharma 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 22, 2021. Such persons may also file a written request for a hearing on the application on or before November 22, 2021.

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: