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:

Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on September 8, 2021, Novitium Pharma LLC, 70 Lake Drive, East Windsor, New Jersey 08520, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Levorphanol	9220	II

The company plans to import the listed controlled substance Levorphanol to develop the manufacturing process for a drug product that will in turn be used to produce a tablet equivalent to the current brand product. No other activity for this drug code 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. 2021–23032 Filed 10–21–21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Service Contract Inventory; Notice of Availability

AGENCY: Justice Management Division, Department of Justice.

ACTION: Notice.

SUMMARY: The Department of Justice is publishing this notice to advise the public of the availability of its FY 2019 Service Contracts Inventory and Inventory Supplement. The inventory includes service contract actions over $25,000 that were awarded in Fiscal Year (FY) 2019. The inventory supplement includes information collected from contractors on the amount invoiced and direct labor hours expended for covered service contracts. The Department of Justice analyzes this data for the purpose of determining whether its contract labor is being used in an effective and appropriate manner and if the mix of federal employees and contractors in the agency is effectively balanced. The inventory and supplement do not include contractor proprietary or sensitive information. The FY 2019 Service Contract Inventory and Inventory Supplements are provided at the following link: *https://www.justice.gov/jmd/service-contractinventory.*

FOR FURTHER INFORMATION CONTACT: Kevin Doss, Office of Acquisition Management, Justice Management Division, U.S. Department of Justice, Washington, DC 20530; Phone: 202–616–3758; Email: *Kevin.Doss@usdoj.gov.*

Authority: Section 743 of Division C of the FY 2010 Consolidated Appropriations Act, Pub. L. 111–117.

Dated: October 18, 2021.

Melody Braswell,

Department Clearance Officer for PRA, U.S. Department of Justice.

[FR Doc. 2021–23014 Filed 10–21–21; 8:45 am]

BILLING CODE 4410–02–P

DEPARTMENT OF LABOR

Occupational Safety and Health Administration

[Docket No. OSHA–2021–0010]

Federal Advisory Council on Occupational Safety and Health

AGENCY: Occupational Safety and Health Administration (OSHA), Labor.

ACTION: Notice of reestablishment of the Federal Advisory Council on Occupational Safety and Health (hereinafter FACOSH or Council) and request for nominations to serve on FACOSH.

SUMMARY: The Occupational Safety and Health Administration (OSHA) hereby announces that the Federal Advisory Council on Occupational Safety and Health (FACOSH) has been reestablished for a two-year period pursuant to the Federal Advisory Committee Act (FACA) and in accordance with the Committee Management Secretariat, General Services Administration. In addition, the Secretary of Labor (Secretary) requests nominations for membership on FACOSH.

DATES: Submit (postmark, send, transmit) nominations for FACOSH membership by November 22, 2021.

ADDRESSES: You may submit nominations and supporting materials by one of the following methods:

Electronically: You may submit nominations, including attachments, electronically into Docket No. OSHA—2021–0010 at *http://www.regulations.gov,* which is the Federal eRulemaking Portal. Follow the online instructions for submissions.

Docket: To read or download comments or other material in the docket, go to *http://www.regulations.gov.* Documents in the docket are listed in the *http://www.regulations.gov* index; however, some information (*e.g.,* copyrighted material) is not publicly available to read or download through the website. All submissions, including copyrighted material, are available for inspection through the OSHA Docket Office. Contact the OSHA Docket Office at (202) 693–2350 (TTY (877) 889–5627) for assistance in locating docket submissions.

Instructions: All submissions must include the agency name and the OSHA docket number for this **Federal Register** notice (OSHA–2021–0010). OSHA will place comments and requests to speak, including personal information, in the public docket, which may be available online. Therefore, OSHA cautions interested parties about submitting personal information such as Social Security numbers and birthdates.

FOR FURTHER INFORMATION CONTACT:

Press inquiries: Mr. Frank Meilinger, Director, OSHA Office of Communications; telephone: (202) 693–1999; email: *meilinger.francis2@dol.gov.*

General information: Mr. Francis Yebesi, Director, OSHA Office of Federal Agency Programs; telephone (202) 693–2122; email *ofap@dol.gov.*

Copies of this **Federal Register** *document:* Electronic copies of this **Federal Register** document are available at *http://www.regulations.gov.* This document, as well as news releases and other relevant information are also available on the OSHA web page at *http://www.osha.gov.*

SUPPLEMENTARY INFORMATION: On September 30, 2021, President Joseph Biden signed Executive Order (E.O.) 14048 continuing or reestablishing certain federal advisory committees, including FACOSH, until September 30, 2023 (86 FR 55465 (10/05/2021)). In response, the Secretary reestablished FACOSH and the Department of Labor (DOL) filed the FACOSH charter on October 14, 2021. FACOSH will terminate on September 30, 2023, unless continued by the President. The FACOSH charter is available to read or download at *https://www.osha.gov/.* In addition, the Secretary invites interested persons to submit nominations for membership on FACOSH.