European customers, meets the standards established by the European Pharmacopeia, administered by the Directorate for the quality of Medicines (EDQM). In order to ensure that its product will meet European specifications, the company seeks to import morphine supplied by EDQM for use as reference standards. 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.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–26172 Filed 11–25–20; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–743]

Bulk Manufacturer 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 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 January 26, 2021. Such persons may also file a written request for a hearing on the application on or before January 26, 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 November 2, 2020, Novitium Pharma LLC, 70 Lake Drive, East Windsor, New Jersey 08520, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substances:

Controlled substance	Drug code	Schedule
Levorphanol	9220	II

The company plans to bulk manufacture the above-controlled substance to support production of the company's Food and Drug Administration approved drug product. No other activity for this drug code is authorized for this registration.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–26171 Filed 11–25–20; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–746]

Importer of Controlled Substances Application: Noramco Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Noramco Inc. 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 December 28, 2020. Such persons may also file a written request for a hearing on the application on or before December 28, 2020.

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 October 23, 2020, Noramco Inc., 500 Swedes Landing Road, Wilmington, Delaware, 19801–4417, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Marihuana	7360	I
Tetrahydrocannabinols	7370	I
Nabilone	7379	II
Phenylacetone	8501	II
Opium, Raw	9600	II
Poppy Straw Concentrate	9670	II
Tapentadol	9780	II

The company plans to import Phenylacetone (8501), and Poppy Straw Concentrate (9670) to bulk manufacture other controlled substances for distribution to its customers. The company plans to import an intermediate form of Tapentadol (9780) to bulk manufacture Tapentadol for distribution to its customers. In reference to drug codes 7360 (Marihuana) and 7370 (Tetrahydrocannabinols), the company plans to import a synthetic cannabidiol and a synthetic Tetrahydrocannabinol. 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.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–26174 Filed 11–25–20; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–742]

Importer of Controlled Substances Application: Lyndra Therapeutics

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Lyndra Therapeutics 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 December 28, 2020. Such persons may also file a written request for a hearing on the application on or before December 28, 2020.

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 October 22, 2020, Lyndra Therapeutics, 65 Grove Street, Suite 301, Watertown, Massachusetts 02472, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Methadone	9250	II

The company plans to develop the formulation and process, and then manufacture the finished oral dosage form for use in preclinical and human clinical trials under a research grant from National Institute on Drug Abuse. 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.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–26175 Filed 11–25–20; 8:45 am]

BILLING CODE P

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[Notice: (20–097)]

NASA Advisory Council; Science Committee; Meeting

AGENCY: National Aeronautics and Space Administration.

ACTION: Notice of revised dates for meeting.

SUMMARY: In accordance with the Federal Advisory Committee Act, as amended, the National Aeronautics and Space Administration (NASA) announces revised dates for the upcoming meeting of the Science Committee of the NASA Advisory Council (NAC).

DATES: Tuesday, December 1, 2020, 1:00 p.m.–5:00 p.m., and Wednesday, December 2, 2020, 1:00–5:15 p.m., Eastern Time.

FOR FURTHER INFORMATION CONTACT: Ms. KarShelia Henderson, Science Mission Directorate, NASA Headquarters, Washington, DC 20546, (202) 358–2355 or *khenderson@nasa.gov.*

SUPPLEMENTARY INFORMATION: The original meeting notice was published in the **Federal Register** on Friday, November 13, 2020 in Vol. 85, No. 220 on page 72703. This meeting will now take place on two days (December 1–2, 2020), rather than on three days (December 1–3, 2020). The Science Committee reports to the NAC. The meeting will be held for the purpose of soliciting, from the scientific community and other persons, scientific and technical information relevant to program planning. This meeting will be open to the public via Webex and telephonically. Webex connectivity information for each day is provided below. For audio, when you join the Webex event, you may use your computer or provide your phone number to receive a call back, otherwise, call the U.S. toll conference number listed for each day. On Tuesday, December 1, the event address for attendees is: *https://nasaenterprise.webex.com/nasaenterprise/onstage/g.php?MTID=ec9f04af53099d097214a64cf178fc2ed.* The event number is 199 056 0375 and the event password is wfSEe8uH5*3. If needed, the U.S. toll conference number is 1–415–527–5035 and access code is 199 056 0375. On Wednesday, December 2, the event address for attendees is: *https://nasaenterprise.webex.com/nasaenterprise/onstage/g.php?MTID=e51f38c7ac92a01577c5f697d7d1b4c5f.* The event number is 199 748 1916 and the event password is EswGXYZ@742. If needed, the U.S. toll conference number is 1–415–527–5035 and access code is 199 748 1916.

The agenda for the meeting includes the following topics:

—Science Mission Directorate (SMD) Missions, Programs and Activities

It is imperative that the meeting be held on these dates due to the scheduling priorities of the key participants.

Patricia Rausch,

Advisory Committee Management Officer, National Aeronautics and Space Administration.

[FR Doc. 2020–26244 Filed 11–25–20; 8:45 am]

BILLING CODE 7510–13–P

NATIONAL LABOR RELATIONS BOARD

Privacy Act of 1974; System of Records

AGENCY: National Labor Relations Board (NLRB).

ACTION: Notice of a new privacy act system of records.

SUMMARY: In accordance with the Privacy Act of 1974, the National Labor Relations Board proposes to issue a National Labor Relations Board system of records notice titled ''NLRB iTrak and Banned Entry List'' (NLRB–34) to support the protection of employees, contractors, and property leased, or occupied, by the National Labor Relations Board. This system of records includes the NLRB's iTrak Incident & Security Management Software System (''iTrak''), which is used to manage information on individuals who have been reported to present a threat or potential threat to NLRB employees, contractors, and property, as well as a Banned Entry List, which is a list of individuals banned from entering NLRB facilities based on information in iTrak. The system allows the National Labor Relations Board to collect and maintain records on the results of law enforcement activities concerning individuals maintaining a presence at or who have access to property leased or occupied by the NLRB and who have been reported to present a threat as described above. The NLRB is issuing this system of records notice in compliance with the Privacy Act of