Controlled substance	Drug code	Schedule
Morphine	9300	II

The company plans to synthesize the above listed control substances for distribution to its customers. In reference to drug codes 7360 (Marihuana) and 7370 (Tetrahydrocannabinol), the company plans to bulk manufacture these drugs as synthetics. No other activity for these drug codes are authorized for this registration.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2021–00356 Filed 1–11–21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–762]

Importer of Controlled Substances Application: S&B Pharma LLC dba Norac Pharma

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: S&B Pharma LLC, dba: Norac Pharma 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 February 11, 2021. Such persons may also file a written request for a hearing on the application on or before February 11, 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 August 25, 2020, S&B Pharma LLC, dba: Norac Pharma, 405 South Motor Avenue, Azusa, California 91702, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
4-Anilino-N-phenethyl-4-piperidine (ANPP).	8333	II
Tapentadol	9780	II

The company plans to import the listed controlled substances in bulk for the manufacture of 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 Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2021–00326 Filed 1–11–21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–764]

Bulk Manufacturer of Controlled Substances Application: Siegfried USA, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Siegfried USA, 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 March 15, 2021. Such persons may also file a written request for a hearing on the application on or before March 15, 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 December 9, 2020, Siegfried USA, LLC, 33 Industrial Park Road, Pennsville, New Jersey 08070–3244, 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
Amphetamine	1100	II
Methylphenidate	1724	II
Amobarbital	2125	II
Pentobarbital	2270	II
Secobarbital	2315	II
Codeine	9050	II
Oxycodone	9143	II
Hydromorphone	9150	II
Hydrocodone	9193	II
Methadone	9250	II
Methadone intermediate	9254	II
Morphine	9300	II
Oripavine	9330	II
Thebaine	9333	II
Opium tincture	9630	II
Oxymorphone	9652	II
Tapentadol	9780	II

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

William T. McDermott,

Assistant Administrator.

[FR Doc. 2021–00354 Filed 1–11–21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–760]

Bulk Manufacturer of Controlled Substances Application: Bulk Manufacturer of Marihuana: Natural Fulfillment LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: The Drug Enforcement Administration (DEA) is providing notice of an application it has received from an entity applying to be registered to manufacture in bulk basic class(es) of controlled substances listed in schedule I. DEA intends to evaluate this and other pending applications according to its regulations governing the program of growing marihuana for scientific and medical research under DEA registration.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefor, may file written comments on or objections to the issuance of the proposed registration on or before March 15, 2021.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW 8701 Morrissette Drive, Springfield, Virginia 22152. To ensure proper handling of comments, please reference Docket No—DEA–760 in all correspondence, including attachments.

SUPPLEMENTARY INFORMATION: The Controlled Substances Act (CSA) prohibits the cultivation and distribution of marihuana except by persons who are registered under the CSA to do so for lawful purposes. In accordance with the purposes specified in 21 CFR 1301.33(a), DEA is providing notice that the entity identified below has applied for registration as a bulk manufacturer of schedule I controlled substances. In response, registered bulk manufacturers of the affected basic class(es), and applicants therefor, may file written comments on or objections of the requested registration, as provided in this notice. This notice does not constitute any evaluation or determination of the merits of the application submitted.

The applicant plans to manufacture bulk active pharmaceutical ingredients (APIs) for product development and distribution to DEA registered researchers. If the application for registration is granted, the registrant would not be authorized to conduct other activity under this registration aside from those coincident activities specifically authorized by DEA regulations. DEA will evaluate the application for registration as a bulk manufacturer for compliance with all applicable laws, treaties, and regulations and to ensure adequate safeguards against diversion are in place.

As this applicant has applied to become registered as a bulk manufacturer of marihuana, the application will be evaluated under the criteria of 21 U.S.C. 823(a). DEA will conduct this evaluation in the manner described in the rule published at 85 FR 82333 on December 18, 2020, and reflected in DEA regulations at 21 CFR part 1318.

In accordance with 21 CFR 1301.33(a), DEA is providing notice that on December 1, 2020, Natural Fulfillment LLC, 5495 North Academy Boulevard, Colorado Springs, Colorado 80918, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substances:

Controlled substance	Drug code	Schedule
Marihuana	7360	I

William T. McDermott,

Assistant Administrator.

[FR Doc. 2021–00327 Filed 1–11–21; 8:45 am]

BILLING CODE P

NATIONAL SCIENCE FOUNDATION

Notice of Open to the Public Meetings of the Networking and Information Technology Research and Development (NITRD) Program

AGENCY: Networking and Information Technology Research and Development (NITRD) National Coordination Office (NCO), National Science Foundation.

ACTION: Notice of public meetings.

SUMMARY: The NITRD Program holds meetings that are open to the public to attend. The Joint Engineering Team (JET) and Middleware And Grid Interagency Coordination (MAGIC) Team provide an opportunity for the public to engage and participate in information sharing with Federal agencies. The JET and MAGIC Teams report to the NITRD Large Scale Networking (LSN) Interagency Working Group (IWG).

DATES: January 2021–December 2021.

FOR FURTHER INFORMATION CONTACT: NITRD NCO at *admin@nitrd.gov.* Individuals who use a telecommunications device for the deaf (TDD) may call the Federal Information Relay Service (FIRS) at 1–800–877–8339 between 8 a.m. and 8 p.m., Eastern time, Monday through Friday.

SUPPLEMENTARY INFORMATION: The Joint Engineering Team (JET), established in 1997, provides an opportunity for information sharing among Federal agencies and non-Federal participants who have an interest in high-performance research and engineering or research and education (REN) networking and networking to support science applications.

The MAGIC Team, established in 2002, provides for information sharing among Federal agencies and non-Federal participants with interests and responsibility for middleware, Grid, and cloud projects; individuals involved in middleware, Grid, and cloud research and infrastructure; individuals involved in implementing or operating Grids and clouds; and users of Grids, clouds and middleware. The JET and MAGIC Team meetings are hosted by the NITRD NCO with WebEx and/or teleconference participation available for each meeting.

Public Meetings Website: The JET and MAGIC Team meetings are scheduled 30 days in advance of the meeting date. Please reference the NITRD Public Meetings web page (*https://www.nitrd.gov/meetings/public/*) for each Team's upcoming meeting dates and times, in addition to the agendas, minutes, and other meeting materials and information.

Public Meetings Mailing Lists: Members of the public may be added to the mailing lists by sending their full name and email address to *jet-signup@nitrd.gov* for JET and *magic-signup@nitrd.gov* for MAGIC, with the subject line: ''Add to JET'' and/or ''Add to MAGIC.'' Meeting notifications and information are shared via the mailing lists.

Public Comments: The government seeks individual input; attendees/participants may provide individual advice only. Members of the public are welcome to submit their comments for JET to *jet-comments@nitrd.gov* and for MAGIC to *magic-comments@nitrd.gov*. Please note that under the provisions of the Federal Advisory Committee Act (FACA), all public comments and/or presentations will be treated as public documents and may be made available to the public via the JET and MAGIC web pages.

Reference Website: NITRD website at: *http://www.nitrd.gov/*.

Submitted by the National Science Foundation in support of the Networking and Information Technology Research and Development (NITRD) National Coordination Office (NCO) on January 6, 2021.

Suzanne H. Plimpton,

Reports Clearance Officer, National Science Foundation.

[FR Doc. 2021–00365 Filed 1–11–21; 8:45 am]

BILLING CODE 7555–01–P