SUMMARY: The Commission instituted the subject five-year review on August 1, 2025 to determine whether revocation of the antidumping duty order on Polyethylene Terephthalate (PET) Sheet from South Korea would be likely to lead to continuation or recurrence of material injury. Due to the lapse in appropriations and Federal Government shutdown, on November 14, 2025, the Department of Commerce (''Commerce'') tolled all deadlines in administrative proceedings by 47 days. Additionally, due to a backlog of documents that were electronically filed via Enforcement and Compliance's Antidumping and Countervailing Duty Centralized Electronic Service System (ACCESS) during the Federal Government shutdown, on November 24, 2025, Commerce tolled all deadlines in administrative proceedings by an additional 21 days. On January 12, 2026, Commerce published notice in the **Federal Register** that it was revoking the order effective January 12, 2026, because no domestic interested party filed a timely notice of intent to participate. Accordingly, the subject review is terminated.

DATES: January 12, 2026.

FOR FURTHER INFORMATION CONTACT: Rachel Devenney (202–205–3172), Office of Investigations, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436. Hearing-impaired individuals are advised that information on this matter can be obtained by contacting the Commission's TDD terminal on 202–205–1810. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at 202–205–2000. General information concerning the Commission may also be obtained by accessing its internet server (*https://www.usitc.gov*). The public record for this investigation may be viewed on the Commission's electronic docket (EDIS) at *https://edis.usitc.gov.*

Authority: This review is being terminated under authority of title VII of the Tariff Act of 1930 and pursuant to section 751(c) of the Tariff Act of 1930 (19 U.S.C. 1675(c)). This notice is published pursuant to section 207.69 of the Commission's rules (19 CFR 207.69).

By order of the Commission.

Issued: January 30, 2026.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2026–02235 Filed 2–3–26; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[Investigation Nos. 701–TA–499–500 and 731–TA–1215–1216, 1221–1223 (Second Review)]

Oil Country Tubular Goods From India, South Korea, Turkey, Ukraine, and Vietnam; Notice of Commission Determination To Conduct Full Five-Year Reviews

AGENCY: United States International Trade Commission.

ACTION: Notice.

SUMMARY: The Commission hereby gives notice that it will proceed with full reviews pursuant to the Tariff Act of 1930 to determine whether revocation of the countervailing duty orders on oil country tubular goods from India and Turkey and the antidumping duty orders on oil country tubular goods from India, South Korea, Turkey, Ukraine, and Vietnam would be likely to lead to continuation or recurrence of material injury within a reasonably foreseeable time. A schedule for the reviews will be established and announced at a later date.

DATES: November 24, 2025.

FOR FURTHER INFORMATION CONTACT: Peter Stebbins (202–205–20239), Office of Investigations, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436. Hearing-impaired persons can obtain information on this matter by contacting the Commission's TDD terminal on 202–205–1810. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at 202–205–2000. General information concerning the Commission may also be obtained by accessing its internet server (*https://www.usitc.gov*). The public record for these reviews may be viewed on the Commission's electronic docket (EDIS) at *https://edis.usitc.gov.*

For further information concerning the conduct of these reviews and rules of general application, consult the Commission's Rules of Practice and Procedure, part 201, subparts A through E (19 CFR part 201), and part 207, subparts A, D, E, and F (19 CFR part 207).

SUPPLEMENTARY INFORMATION: On November 24, 2025, the Commission determined that it should proceed to full reviews in the subject five-year reviews pursuant to section 751(c) of the Tariff Act of 1930 (19 U.S.C. 1675(c)).[1] The Commission found that both the domestic and respondent interested party group responses from Ukraine to its notice of institution (90 FR 28768, July 1, 2025) were adequate, and determined to conduct a full review of the order on imports from Ukraine. The Commission also found that the respondent interested party group responses from India, Turkey, South Korea, and Vietnam were inadequate but determined to conduct full reviews of the orders on imports from those countries in order to promote administrative efficiency in light of its determinations to conduct full reviews of the orders with respect to Ukraine. A record of the Commissioners' votes will be available from the Office of the Secretary and at the Commission's website.

Authority: These reviews are being conducted under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to § 207.62 of the Commission's rules.

By order of the Commission.

Issued: January 30, 2026.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2026–02210 Filed 2–3–26; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1658]

Bulk Manufacturer 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 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 submit electronic comments on or objections to the issuance of the proposed registration on or before April 6, 2026. Such persons may also file a written request for a hearing on the application on or before April 6, 2026.

ADDRESSES: The Drug Enforcement Administration requires that all

[1] Due to the lapse in appropriations and ensuing cessation of Commission operations, the Commission tolled its schedule for this proceeding.

comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to *https://www.regulations.gov* and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on *https://www.regulations.gov.* If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on January 13, 2026, S & B Pharma LLC DBA Norac Pharma, 405 South Motor Avenue, Azusa, California 91702, applied to be registered as a 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 above listed controlled substances for internal research and for development purposes as part of the process in seeking Food and Drug Administration approval prior to distribution to customers. No other activities for these drug codes are authorized for this registration.

Thomas Prevoznik,

Deputy Assistant Administrator.

[FR Doc. 2026–02232 Filed 2–3–26; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1655]

Bulk Manufacturer of Controlled Substances Application: Sterling Pharma USA LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Sterling Pharma USA 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 submit electronic comments on or objections to the issuance of the proposed registration on or before April 6, 2026. Such persons may also file a written request for a hearing on the application on or before April 6, 2026.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to *https://www.regulations.gov* and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on *https://www.regulations.gov.* If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on January 9, 2026, Sterling Pharma USA LLC, 1001 Sheldon Drive, Suite 101, Cary, North Carolina 27513–2078, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Tetrahydrocannabinols	7370	I
5-Methoxy-N-N-dimethyltryptamine.	7431	I
Dimethyltryptamine	7435	I
Psilocybin	7437	I
Psilocyn	7438	I

The company plans to bulk manufacture the listed controlled substances to support internal research and for sale to its customers for pre-clinical trial studies. No other activities for these drug codes are authorized for this registration.

Thomas Prevoznik,

Deputy Assistant Administrator.

[FR Doc. 2026–02230 Filed 2–3–26; 8:45 am]

BILLING CODE 4410–09–P

NUCLEAR REGULATORY COMMISSION

[NRC–2025–2161]

Duke Energy Carolinas, LLC; Belews Creek; Early Site Permit Application

AGENCY: Nuclear Regulatory Commission.

ACTION: Notice; receipt.

SUMMARY: The U.S. Nuclear Regulatory Commission (NRC) is providing public notice each week, for four consecutive weeks, of receipt and availability of an application for an early site permit (ESP) from Duke Energy Carolinas, LLC, for the Belews Creek site located in Stokes County, North Carolina.

DATES: February 4, 2026.

ADDRESSES: Please refer to Docket ID NRC–2025–2161 when contacting the NRC about the availability of information regarding this document. You may obtain publicly available information related to this document using any of the following methods:

- *Federal Rulemaking Website:* Go to *https://www.regulations.gov* and search for Docket ID NRC–2025–2161. Address questions about Docket IDs in *Regulations.gov* to Bridget Curran; telephone: 301–415–1003; email: *Bridget.Curran@nrc.gov.* For technical questions, contact the individual(s) listed in the For **FURTHER INFORMATION CONTACT** section of this document.
- *NRC's Agencywide Documents Access and Management System (ADAMS):* You may obtain publicly available documents online in the ADAMS Public Documents collection at *https://www.nrc.gov/reading-rm/adams.html.* To begin the search, select ''Begin ADAMS Public Search.'' For problems with ADAMS, please contact the NRC's Public Document Room (PDR) reference staff at 1–800–397–4209, at 301–415–4737, or by email to *PDR.Resource@nrc.gov.* The Belews Creek Early Site Permit Application package is available in ADAMS under Accession No. ML25364A004.
- *NRC's PDR:* The PDR, where you may examine and order copies of publicly available documents, is open by appointment. To make an appointment to visit the PDR, please send an email to *PDR.Resource@nrc.gov* or call 1–800–397–4209 or 301–415–4737, between 8 a.m. and 4 p.m. eastern time (ET), Monday through Friday, except Federal holidays.

FOR FURTHER INFORMATION CONTACT: Emmanuel Sayoc, Office of Nuclear Reactor Regulation, U.S. Nuclear Regulatory Commission, Washington, DC 20555–0001; telephone: 301–415–4084; email: *Emmanuel.Sayoc@nrc.gov.*