

requirements for other Midwest states that limit the volatility of E10 in those states, as provided in 40 CFR 1090.215(b)(3).

Aaron Szabo,

Assistant Administrator, Office of Air and Radiation.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS-10791]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by April 6, 2026.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to <http://www.regulations.gov>. Follow the instructions for "Comment or Submission" or "More Search Options" to find the information collection

document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier: ____ / OMB Control Number: ____, Room C4-26-05, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, please access the CMS PRA website by copying and pasting the following web address into your web browser: <https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing>.

FOR FURTHER INFORMATION CONTACT: William N. Parham at (410) 786-4669.

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

Under the PRA (44 U.S.C. 3501-3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term "collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

Information Collections

1. *Type of Information Collection Request:* Reinstatement without change of a previously approved collection; *Title of Information Collection:* Requirements Related to Surprise Billin; Part II; *Use:* The collection of information is associated with the October 7, 2021 (86 FR 55980) interim final rules. The collection has two components:

A. *Good Faith Estimates.* Providers and facilities must inform uninsured (or

self-pay) individuals of their right to receive a good faith estimate (GFE) of expected charges for items and services. They must also furnish a good faith estimate of expected charges to uninsured (or self-pay) individuals for scheduled items and services and upon request, which provides uninsured (or self-pay) individuals information about health care pricing before receiving care. This information would allow uninsured (or self-pay) individuals to evaluate options for receiving health care and make cost-conscious health care purchasing decisions and reduces surprises regarding individuals' health care costs for items and services. Additionally, uninsured (or self-pay) individuals need a good faith estimate to initiate the patient-provider dispute resolution process.

B. *Certification and Recertification of SDR Entities.* HHS requests information from entities seeking to be certified or recertified as an SDR entity. This information is used to assess whether or not the entity satisfies the requirements for certification. Entities must submit information on their organizational structure, policies and procedures, staff qualifications, conflict-of-interest safeguards, and operational capacity, along with attestations of compliance with applicable standards. This information allows HHS to determine the entity's eligibility and capability to perform SDR functions effectively and impartially. *Form Number:* CMS-10791 (OMB control number: 0938-1433); *Frequency:* Annually; *Affected Public:* Private sector (Business or other for-profits and Not-for-profit institutions); *Number of Respondents:* 511,749; *Total Annual Responses:* 5,248,414; *Total Annual Hours:* 3,498,944. (For policy questions regarding this collection contact Daniel Kidane at daniel.kidane@cms.hhs.gov.)

William N. Parham, III,

Director, Division of Information Collections and Regulatory Impacts, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2026-02215 Filed 2-3-26; 8:45 am]

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INTERNATIONAL TRADE COMMISSION

[Investigation No. 731-TA-1455 (Review)]

Polyethylene Terephthalate (PET) Sheet From South Korea; Termination of Five-Year Review

AGENCY: United States International Trade Commission.

ACTION: Notice.

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]

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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.

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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.