

(iii) identify like or directly competitive articles that complainant, its licensees, or third parties make in the United States which could replace the subject articles if they were to be excluded;

(iv) indicate whether complainant, complainant's licensees, and/or third party suppliers have the capacity to replace the volume of articles potentially subject to the requested exclusion order and/or a cease and desist order within a commercially reasonable time; and

(v) explain how the requested remedial orders would impact United States consumers.

Written submissions on the public interest must be filed no later than by close of business, eight calendar days after the date of publication of this notice in the **Federal Register**. There will be further opportunities for comment on the public interest after the issuance of any final initial determination in this investigation. Any written submissions on other issues must also be filed by no later than the close of business (5:15 p.m.),¹ eight calendar days after publication of this notice in the **Federal Register**.

Complainant(s) may file replies to any written submissions no later than three calendar days after the date on which the written submissions were due. Notwithstanding § 201.14(a) of the Commission's Rules of Practice and Procedure, the deadlines for written submissions and replies shall be counted by calendar days starting with the first calendar day following the **Federal Register** publication date or the due date for written submissions, respectively. The provisions of § 201.14(a) regarding the last day of the period still apply concerning the due date of any filings. No other submissions will be accepted, unless requested by the Commission. Any submissions and replies filed in response to this Notice are limited to five (5) pages in length, inclusive of attachments.

Persons filing written submissions must file the original document electronically on or before the deadlines stated above. Submissions should refer to the docket number ("Docket No. 3939") in a prominent place on the cover page and/or the first page. (See Handbook for Electronic Filing Procedures, Electronic Filing Procedures²). Please note the Secretary's Office will accept only

electronic filings unless an exemption is granted. Filings must be made through the Commission's Electronic Document Information System (EDIS, <https://edis.usitc.gov>.) Persons with questions regarding filing should contact the Secretary at EDIS3Help@usitc.gov.

Any person desiring to submit a document to the Commission in confidence must request confidential treatment. All such requests should be directed to the Secretary to the Commission and must include a full statement of the reasons why the Commission should grant such treatment. See 19 CFR 201.6. Documents for which confidential treatment by the Commission is properly sought will be treated accordingly. All information, including confidential business information and documents for which confidential treatment is properly sought, submitted to the Commission for purposes of this Investigation may be disclosed to and used: (i) by the Commission, its employees and Offices, and contract personnel (a) for developing or maintaining the records of this or a related proceeding, or (b) in internal investigations, audits, reviews, and evaluations relating to the programs, personnel, and operations of the Commission including under 5 U.S.C. Appendix 3; or (ii) by U.S. government employees and contract personnel,³ solely for cybersecurity purposes. All nonconfidential written submissions will be available for public inspection at the Office of the Secretary and on EDIS.⁴

This action is taken under the authority of section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and of §§ 201.10 and 210.8(c) of the Commission's Rules of Practice and Procedure (19 CFR 201.10, 210.8(c)).

By order of the Commission.

Issued: September 28, 2026.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2026-20081 Filed 9-30-26; 8:45 am]

BILLING CODE 7020-02-P

DEPARTMENT OF JUSTICE

[OMB Number 1117-0056]

Agency Information Collection Activities; Proposed eCollection Comments Requested; Proposed New Collection Request; Title—Suspicious Order of Controlled Substances

AGENCY: Drug Enforcement Administration, Department of Justice.

ACTION: 30-day Notice.

SUMMARY: The Department of Justice (DOJ), Drug Enforcement Administration (DEA), will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. This proposed information collection was previously published in the **Federal Register** on July 29, 2026, at 91 FR 47856, allowing for a 60-day comment period.

DATES: Comments are encouraged and will be accepted for 30 days until November 2, 2026.

FOR FURTHER INFORMATION CONTACT: If you have comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact Heather Achbach, Regulatory Drafting and Policy Support Section, Drug Enforcement Administration; Mailing Address: 8701 Morrisette Drive, Springfield, Virginia 22152; Telephone: (571) 776-3822, email: DEA.PRA@dea.gov.

SUPPLEMENTARY INFORMATION: Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

- Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
- Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- Enhance the quality, utility, and clarity of the information to be collected; and/or
- Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic,

¹ See 19 CFR 201.3(c).

² Handbook for Electronic Filing Procedures: https://www.usitc.gov/secretary/documents/handbook_on_filing_procedures.pdf.

³ All contract personnel will sign appropriate nondisclosure agreements.

⁴ Electronic Document Information System (EDIS): <https://edis.usitc.gov>.

mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

Written comments and recommendations for this information collection should be submitted within 30 days of the publication of this notice on the following website www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function and entering either the title of the information collection or the OMB Control Number 1117–0003. This information collection request may be viewed at www.reginfo.gov. Follow the instructions to view Department of Justice, information collections currently under review by OMB. If you submit public comments, please send a copy to *Heather Achbach at DEA.PRA@dea.gov*.

DOJ seeks PRA authorization for this information collection for three (3) years. OMB authorization for an ICR cannot be for more than three (3) years without renewal. The DOJ notes that information collection requirements submitted to the OMB for existing ICRs receive a month-to-month extension while they undergo review.

Abstract: This collection is a centralized database for the reporting of all suspicious orders of controlled substances. Public comment was also solicited in the “Suspicious Orders of Controlled Substances” notice of proposed rulemaking (NPRM), published in the **Federal Register** at 85 FR 69282, on November 2, 2020. The NPRM had a 60-day comment period. No comments on PRA were received.

Overview of This Information Collection

- 1. *Type of Information Collection:* Proposed new collection request.

- 2. *Title of the Form/Collection:* Suspicious Orders of Controlled Substances.
- 3. *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection:* There will be no form number. The applicable component within the Department of Justice is the Drug Enforcement Administration, Diversion Control Division.
- 4. *Affected public who will be asked or required to respond, as well as a brief abstract:* Affected public (Primary): Business or other for-profit.
- 5. *Total Estimated Number of Respondents:* 178.
- 6. *Estimated Time per Respondent:* The time per response is 0.16669 minutes to complete the report.
- 7. *Frequency:* 4,154.096 per year.
- 8. *Total Estimated Annual Time Burden:* 12,324 hours.
- 9. *Total Estimated Annual Other Costs Burden:* \$0.

TOTAL BURDEN HOURS

Activity	Number of respondents	Frequency	Total annual responses	Time per response (hours)	Total annual burden (hours)
Reporting of Suspicious Orders	178	4,154.096	739,429	0.01666691	12,324
Total	178	N/A	739,429	0.01666691	12,324

If additional information is required, contact: Darwin Arceo, Department Clearance Officer, Enterprise Portfolio Management, Justice Management Division, United States Department of Justice, Two Constitution Square, 145 N Street NE, 4W–218, Washington, DC 20530.

Dated: September 29, 2026.

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

[FR Doc. 2026–20114 Filed 9–30–26; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

[OMB Number 1117–0061]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Revision of a Previously Approved Collection; Title—Recordkeeping for the Transfer of Electronic Prescriptions in Schedules III–V Between Pharmacies

AGENCY: Drug Enforcement Administration, Department of Justice.
ACTION: 30-day Notice.

SUMMARY: The Drug Enforcement Administration, Department of Justice (DOJ), will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The proposed information collection was previously published in the **Federal Register**, volume 91 FR 46484, on July 23, 2026, allowing a 60-day comment period.

DATES: Comments are encouraged and will be accepted for 30 days until November 2, 2026.

FOR FURTHER INFORMATION CONTACT: If you have comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact: Heather E. Achbach, Regulatory Drafting and Policy Support Section, Drug Enforcement Administration; Mailing Address: 8701 Morrisette Drive, Springfield, Virginia 22152; Telephone: (571) 776–3882; Email: DEA.PRA@dea.gov.

SUPPLEMENTARY INFORMATION:

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

- Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
- Evaluate the accuracy of the agency’s estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- Enhance the quality, utility, and clarity of the information to be collected; and/or
- Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

Written comments and recommendations for this information collection should be submitted within