(St. Louis, Missouri).[1] On July 17, 2026, petitioners withdrew these petitions at both the ITC and the Department of Commerce.[2] Therefore these investigations are hereby terminated.

Authority: These investigations are being terminated under authority of title VII of the Tariff Act of 1930 and pursuant to section 207.40(a) of the Commission's Rules of Practice and Procedure (19 CFR 207.40(a)). This notice is published pursuant to section 201.10 of the Commission's rules (19 CFR 201.10).

By order of the Commission.

Issued: July 24, 2026.

Sharon Bellamy,

Supervisory Hearings and Information Officer.

[FR Doc. 2026–15235 Filed 7–28–26; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

[OMB Number 1117–0056]

Agency Information Collection Activities; Proposed eCollection, eComments Requested; Proposed New Collection Request; Title—Suspicious Orders of Controlled Substances

AGENCY: Drug Enforcement Administration, Department of Justice.

ACTION: 60-Day notice.

SUMMARY: The Drug Enforcement Administration (DEA), 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.

DATES: Comments are encouraged and will be accepted for 60 days until September 28, 2026.

FOR FURTHER INFORMATION CONTACT:

If you have additional 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 Morrissette 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 Bureau of Justice Statistics, 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;

—Evaluate whether and if so, how the quality, utility, and clarity of the information to be collected can be enhanced; and

—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.

Abstract: Section 307 of the Controlled Substances Act (21 U.S.C. 827) requires controlled substance manufacturers and distributors to make periodic reports to DEA regarding the sale, delivery, and other disposal of certain controlled substances. These reports help ensure a closed system of distribution for controlled substances, and are used to comply with international treaty obligations.

Overview of This Information Collection

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

2. *The Title of the Form/Collection:* Suspicious Orders of Controlled Substances.

3. *The agency form number, if any, and the applicable component of the Department 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 a brief abstract:* Affected Public: Business or other for-profit.

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.

5. *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* The DEA estimates that 178 registrants will participate in this information collection. The time per response is 0.16669 minutes to complete the report.

6. *An estimate of the total annual burden (in hours) associated with the collection:* DEA estimates that this collection takes 12,324 annual burden hours.

7. *An estimate of the total annual cost burden associated with the collection, if applicable:* $0.

TOTAL BURDEN HOURS

Activity	Number of respondents	Frequency	Total annual responses	Time per response (hours)	Total annual burden (hours)	* Hourly rate	Monetized value of respondent time
Reporting of Suspicious Orders	178	4,154.096	739,429	0.01666691	12,324	60.82	749,546
Total	178	N/A	739,429	0.01666691	12,324	$60.82	$ 749,546

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

[1] 91 FR 41085 (July 6, 2026).

[2] EDIS Doc. 888337.

Dated: July 27, 2026.

Darwin Arceo,

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

[FR Doc. 2026–15317 Filed 7–28–26; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

[OMB Number 1117–0043]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Title—Medical History and Examination

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.

DATES: Comments are encouraged and will be accepted for 30 days until August 28, 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: Kannessia Jordan, Section Chief, Office of Compliance, Policy Administration Section, 700 Army Navy Drive, Arlington, VA 22202, telephone: 571–776–2262, email: *Kannessia.S.Jordan@DEA.gov.*

SUPPLEMENTARY INFORMATION: The proposed information collection was previously published in the **Federal Register** on date, May 28, 2026, 91 FR 31746, allowing a 60-day comment period. Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged.

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 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 #1117–0043. 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. Please provide a copy of your comments POC Kannessia Jordan, *kannessia.s.jordan@dea.gov,* (517) 776–2262 and reference OMB #1117–0043 in the subject line of your comments.

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.

Overview of This Information Collection

1. *Type of Information Collection:* Existing collection without an OMB number.

2. *Title of the Form/Collection:* Medical History and Examination.

3. *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection:* DEA–325, Drug Enforcement Administration.

4. *Affected public who will be asked or required to respond, as well as a brief abstract:* Affected Public: Individuals.

Abstract: The information collected is the basis for determining medical eligibility of applicants for duty in law enforcement and certain law enforcement support positions with the Drug Enforcement Administration. This information is needed to determine the medical qualifications of applicants based upon their current and past medical history. The information obtained on the DEA–325 ensures the applicant is physically and medically capable of performing the essential duties of the position efficiently and without hazard to themselves or to others. The DEA–325 is needed as part of the required medical examination to assist physicians in making determinations as to medical clearance of applicants for positions within the DEA that are subject to medical standards, and verifies disqualifying medical condition(s) documented on the medical history and examination form.

5. O*bligation to Respond:* Required to obtain employment is positions subject to medical standards, GS–1811 (Special Agents), GS–1801 (Diversion Investigators) and GS–1320 (Forensic Chemists).

6. *Total Estimated Number of Respondents:* 1,800.

7. *Estimated Time per Respondent:* 4 hours.

8. *Frequency:* Once prior to employment.

9. *Total Estimated Annual Time Burden:* 7,200 hours.

10. *Total Estimated Annual Other Costs Burden:* $0 [This is captured in #7 of the 60 day notice, in ROCIS and #13 of Supporting Statement A].

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: July 27, 2026.

Darwin Arceo,

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

[FR Doc. 2026–15305 Filed 7–28–26; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

[OMB Number 1140–0020]

Agency Information Collection Activities; Proposed Collection; Comments Requested; Firearms Transaction Record—ATF Form 5300.9 and 5300.9A (''Form 4473'')

AGENCY: Bureau of Alcohol, Tobacco, Firearms, and Explosives; Department of Justice.

ACTION: Interim notice.

SUMMARY: The Department of Justice (DOJ), Bureau of Alcohol, Tobacco,