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: September 18, 2026.

Sharon Bellamy,

Supervisory Hearings and Information Officer.

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DEPARTMENT OF JUSTICE

[OMB Number 1117–0055]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Extension of a Previously Approved Collection; Title—Procurement Quota Certification and Recordkeeping Requirements

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 November 23, 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) 776–3882; Email: *Heather.E.Achbach@dea.gov* or *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: Pursuant to 21 U.S.C. 826, 21 Cprocure that, and 1315.32(h), any person issued procurement quota for a quantity of a basic class of controlled substances listed in Schedules I or II, and ephedrine, pseudoephedrine, phenylpropanolamine during the current calendar year must, at or before the time of placing an order with a registrant, certify in writing to the registrant from whom they wish to procure that the quantity of such basic class and list I chemical ordered does not exceed the person's unused and available procurement quota of such basic class and chemical for the current calendar year. Registrants shall not fill an order from a person required to apply for a procurement quota unless the order is accompanied by such a certification.

Overview of This Information Collection

1. *Type of Information Collection:* Extension of a currently approved collection.

2. *Title of the Form/Collection:* Procurement Quota Certification and Recordkeeping Requirements.

3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* There is no form for this collection. 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 the obligation to respond:* The affected public is business or other for-profit. The obligation to respond is mandatory to respond to procure any quantity of a basic class of controlled substances listed in Schedules I or II, and ephedrine, pseudoephedrine, phenylpropanolamine.

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 650 registrants participate in this information collection. The time per response is 15 minutes.

6. *An estimate of the total annual burden (in hours) associated with the collection:* DEA estimates that this collection takes 813 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)
Procurement Quota Certification and Recordkeeping	650	5	3,250	0.25	813
Unduplicated Totals	650	5	3,250	0.25	813

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.

Dated: September 21, 2026.

Darwin Arceo,

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

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