

DEPARTMENT OF JUSTICE
[OMB Number 1117-0047]

Agency Information Collection Activities; Proposed eCollection, eComments Requested; Extension Without Change of a Previously Approved Collection; Title—Application for Import Quota for Ephedrine, Pseudoephedrine, and Phenylpropanolamine (DEA Form 488)

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 Morrisette 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. 952 and 21 CFR 1315.34, any person who desires to import the List I chemicals Ephedrine, Pseudoephedrine, or Phenylpropanolamine during the next calendar year must apply on DEA Form

488 for an import quota for each such List I chemical.

Overview of This Information Collection

1. *Type of Information Collection:* Extension of a currently approved collection.
2. *The Title of the Form/Collection:* Application for Import Quota for Ephedrine, Pseudoephedrine, and Phenylpropanolamine.
3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* DEA Form 488. 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:* Affected Public (Primary): Business or other for-profit. *Affected Public (Other):* None.
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 53 registrants participate in this information collection. The time per response is 30 minutes.
6. *An estimate of the total annual burden (in hours) associated with the collection:* DEA estimates that this collection takes 72 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)
DEA Form 488	53	2.69811321	143	0.5	72
Unduplicated Totals	53	2.69811321	143	0.5	72

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.
[FR Doc. 2026-19433 Filed 9-22-26; 8:45 am]

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DEPARTMENT OF JUSTICE
[OMB Number 1117-0008]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Extension of a Previously Approved Collection; Title—Application for Procurement Quota for Controlled Substance and for Ephedrine, Pseudoephedrine, and Phenylpropanolamine (DEA Form 250)

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 Morrisette 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 and 21 CFR 1303.15(b) and 1315.32, any person who desires to use, during the next calendar year, any basic class of controlled substances listed in schedules I or II, or the List I chemicals ephedrine, pseudoephedrine, or phenylpropanolamine for purposes of manufacturing must apply on DEA Form 250 for a procurement quota for such class or List I chemical.

Overview of This Information Collection

1. *Type of Information Collection:* Extension of a currently approved collection.
2. *Title of the Form/Collection:* Application for Procurement Quota for Controlled Substance and for

Ephedrine, Pseudoephedrine, and Phenylpropanolamine.

3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* The form numbers are DEA Forms 250. 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 receive Procurement Quota.

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 262 registrants participate in this information collection. The time per response is 30 minutes.

6. *An estimate of the total annual burden (in hours) associated with the collection:* DEA estimates that this collection takes 1,423 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)
DEA Form 250	262	10.8587786	2,845	0.5	1,423
Unduplicated Totals	262	10.8587786	2,845	0.5	1,423

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.

[FR Doc. 2026-19434 Filed 9-22-26; 8:45 am]

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DEPARTMENT OF JUSTICE

[OMB Number 1117-0006]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Extension Without Change of a Previously Approved Collection; Title—Application for Individual Manufacturing Quota for a Basic Class of Controlled Substance and for Ephedrine, Pseudoephedrine, and Phenylpropanolamine (DEA Form 189)

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 Morrisette 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