

including antidumping or countervailing duties). If you are a trade/business association, provide the information, on an aggregate basis, for the firms which are members of your association.

(a) Production (quantity) and, if known, an estimate of the percentage of total production of *Subject Merchandise* in the *Subject Country* accounted for by your firm's(s') production;

(b) Capacity (quantity) of your firm(s) to produce the *Subject Merchandise* in the *Subject Country* (that is, the level of production that your establishment(s) could reasonably have expected to attain during the year, assuming normal operating conditions (using equipment and machinery in place and ready to operate), normal operating levels (hours per week/weeks per year), time for downtime, maintenance, repair, and cleanup, and a typical or representative product mix); and

(c) the quantity and value of your firm's(s') exports to the United States of *Subject Merchandise* and, if known, an estimate of the percentage of total exports to the United States of *Subject Merchandise* from the *Subject Country* accounted for by your firm's(s') exports.

(12) Identify significant changes, if any, in the supply and demand conditions or business cycle for the *Domestic Like Product* that have occurred in the United States or in the market for the *Subject Merchandise* in the *Subject Country* since the *Order Dates*, and significant changes, if any, that are likely to occur within a reasonably foreseeable time. Supply conditions to consider include technology; production methods; development efforts; ability to increase production (including the shift of production facilities used for other products and the use, cost, or availability of major inputs into production); and factors related to the ability to shift supply among different national markets (including barriers to importation in foreign markets or changes in market demand abroad). Demand conditions to consider include end uses and applications; the existence and availability of substitute products; and the level of competition among the *Domestic Like Product* produced in the United States, *Subject Merchandise* produced in the *Subject Country*, and such merchandise from other countries.

(13) (OPTIONAL) A statement of whether you agree with the above definitions of the *Domestic Like Product* and *Domestic Industry*; if you disagree with either or both of these definitions, please explain why and provide alternative definitions.

**Authority:** This proceeding is being conducted under authority of Title VII of the Tariff Act of 1930; this notice is published pursuant to § 207.61 of the Commission's rules.

By order of the Commission.

Issued: December 21, 2023.

**Lisa Barton,**

*Secretary to the Commission.*

[FR Doc. 2023–28536 Filed 12–29–23; 8:45 am]

**BILLING CODE 7020–02–P**

## JUDICIAL CONFERENCE OF THE UNITED STATES

### Advisory Committee on Appellate Rules; Hearing of the Judicial Conference

**AGENCY:** Judicial Conference of the United States.

**ACTION:** Advisory Committee on Appellate Rules; notice of cancellation of open hearing.

**SUMMARY:** The following public hearing on proposed amendments to the Federal Rules of Appellate Procedure has been canceled: Appellate Rules Hearing on January 24, 2024.

**DATES:** January 24, 2024.

**FOR FURTHER INFORMATION CONTACT:** H. Thomas Byron III, Esq., Chief Counsel, Rules Committee Staff, Administrative Office of the U.S. Courts, Thurgood Marshall Federal Judiciary Building, One Columbus Circle NE, Suite 7–300, Washington, DC 20544, Phone (202) 502–1820, *RulesCommittee\_Secretary@ao.uscourts.gov*.

**SUPPLEMENTARY INFORMATION:** The announcement for this hearing was previously published in the **Federal Register** on August 9, 2023 at 88 FR 53917.

(Authority: 28 U.S.C. 2073.)

Dated: December 27, 2023.

**Shelly L. Cox,**

*Management Analyst, Rules Committee Staff.*

[FR Doc. 2023–28826 Filed 12–29–23; 8:45 am]

**BILLING CODE 2210–55–P**

## DEPARTMENT OF JUSTICE

[OMB Number 1117–0007]

### Agency Information Collection Activities; Proposed eCollection eComments Requested; Registrant Record of Controlled Substances Destroyed

**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. This proposed information collection was previously published in the **Federal Register** on October 26, 2023, allowing for a 60-day comment period.

**DATES:** Comments are encouraged and will be accepted for 30 days until February 1, 2023.

**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 Scott A. Brinks, Regulatory Drafting and Policy Support Section, Drug Enforcement Administration; Mailing Address: 8701 Morrisette Drive, Springfield, Virginia 22152; Telephone: (571) 362–3261; email: *DPW@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 30 days of the publication of this notice on the following website [www.reginfo.gov/public/do/PRAMain](http://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 1117–0007. This information collection request may be viewed at [www.reginfo.gov](http://www.reginfo.gov). Follow the instructions to view Department of Justice, information collections currently under review by OMB.

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: Extension of a currently approved collection.

2. Title of the Form/Collection: Registrant Record of Controlled Substances Destroyed.

3. Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection: DEA Form 41. 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):* Private Sector—business or other for-profit.

*Affected public (Other):* Not-for-profit institutions; Federal, State, local, and tribal governments.

*Abstract:* In accordance with the Controlled Substance Act (CSA), every DEA registrant must make a biennial inventory and maintain, on a current basis, a complete and accurate record of each controlled substance manufactured, received, sold, delivered, or otherwise disposed of. 21 U.S.C. 827 and 958. These records must be maintained separately from all other records of the registrant or, alternatively, in the case of non-narcotic controlled substances, be in such form that required information is readily retrievable from the ordinary business records of the registrant per 21 U.S.C. 827(b)(2). The records must be kept and be available for at least two years for inspection and copying by officers or employees of the United States authorized by the Attorney General. 21 U.S.C. 827(b)(3). The records must be in accordance with and contain such relevant information as may be required by regulations promulgated by DEA. 21

U.S.C. 827(b)(1). These record requirements help to deter and detect diversion of controlled substances and ensure that registrants remain accountable for all controlled substances within their possession and/or control.

5. *Obligation to Respond:* Mandatory per 21 CFR 1314.

6. *Total Estimated Number of Respondents:* 92,832.

7. *Estimated Time per Respondent:* 30 minutes for DEA Form 41.

8. *Frequency:* DEA Form 41 is 1 per year.

9. *Total Estimated Annual Time Burden:* 46,416 hours.

10. *Total Estimated Annual Other Costs Burden:* \$0.

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

Dated: December 27, 2023.

**Darwin Arceo,**

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

[FR Doc. 2023–28816 Filed 12–29–23; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

[OMB Number 1117–0046]

### Agency Information Collection Activities; Proposed eCollection eComments Requested; Self-Certification, Training, and Logbooks for Regulated Sellers and Mail-Order Distributors of Scheduled Listed Chemicals Products

**AGENCY:** Drug Enforcement Administration, Department of Justice.

**ACTION:** 30-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. This proposed information collection was previously published in the **Federal Register** on October 26, 2023, allowing for a 60-day comment period.

**DATES:** Comments are encouraged and will be accepted for 30 days until February 1, 2023.

**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 Scott A. Brinks, Regulatory Drafting and Policy Support Section, Drug Enforcement Administration; Mailing Address: 8701 Morrisette Drive, Springfield, Virginia 22152; Telephone: (571) 362–3261, email: [DPW@dea.gov](mailto:DPW@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 30 days of the publication of this notice on the following website [www.reginfo.gov/public/do/PRAMain](http://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 1117–0046. This information collection request may be viewed at [www.reginfo.gov](http://www.reginfo.gov). Follow the instructions to view Department of Justice, information collections currently under review by OMB.

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.