

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 1117–0010. 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 a comment, 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. DOJ notes that information collection requirements

submitted to the OMB for existing ICRs receive a month-to-month extension while they undergo review.

Abstract: The Controlled Substances Act (CSA) (21 U.S.C. 801–971) establishes a closed system of distribution for controlled substances. One tool that helps to maintain the closed system of distribution is the CSA provision that states it “shall be unlawful for any person to distribute a controlled substance in schedules I or II to another except in pursuance of a written order of the person to whom such substance is distributed, made on a form to be issued by the Attorney General in blank in accordance with subsection (d) of this section.” 21 U.S.C. 828(a).

Overview of This Information Collection

1. *Type of Information Collection:* Reinstatement of a previously approved collection.

2. *Title of the Form/Collection:* U.S. Official Order Forms for Schedules I and II Controlled Substances.

3. *Agency form number, if any, and the applicable component of the*

Department of Justice sponsoring the collection: DEA Form: 222. The applicable component within the Department of Justice is the Drug Enforcement Administration Office of Diversion Control.

4. *Affected public who will be asked or required to respond, as well as a brief abstract:* Affected Public: (Primary): Businesses or other for-profit and not-for-profit institutions; Federal, State, local, and tribal governments.

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 114,989 registrants participate in this information collection. The time per response is 6.19 minutes for the DEA–222.

6. *Estimated Time per Respondent:* DEA estimates that this collection takes 1,624,529 annual burden hours.

7. *Frequency:* As needed, 136.90191/annual.

8. *Total Estimated Annual Time Burden:* 1,624,529 annual burden 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)
DEA–222 (paper)	114,989	2.916644201	335,382	0.25	83,846
DEA–222/CSOS (online)		133.9853	15,406,832	0.1	1,539,846
Unduplicated Totals	114,989	136.90191	15,742,214	0.1031957	1,624,529

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. 2026–20113 Filed 9–30–26; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

[OMB Number 1117–0057]

Agency Information Collection Activities; Proposed eCollection Activities; Proposed eComments Requested; Revision of a Previously Approved Collection; Title—Recordkeeping for Partial Fills of Prescriptions

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. The proposed information collection was previously published in the **Federal Register**, volume 91 FR 46485, 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 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 1117-0057. 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 send a copy of your comments to Heath 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.

Overview of This Information Collection

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

2. *Title of the Form/Collection:* Recordkeeping Requirements for Partial Fills of Prescriptions for Schedule II Controlled Substances.

3. *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection:* No form number is associated with 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:* Affected Public: Business or other for-profit.

5. *Total Estimated Number of Respondents:* 66,088 registrants.

6. *Estimated Time per Respondent:* less than a minute.

7. *Frequency:* 147.8874 as needed, annually.

8. *Total Estimated Annual Time Burden:* 27,149 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)
<i>Pharmacy record of partial fill</i>	66,088	147.8874	9,773,582	0.002778	27,149
Unduplicated Totals	66,088	0	9,773,582	0.002778	27,149

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 2053.

Dated: September 29, 2026.

Darwin Arceo,

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

[FR Doc. 2026-20144 Filed 9-30-8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Notice of Lodging of Proposed Third Consent Decree Modification Under the Comprehensive Environmental Response, Compensation, and Liability Act

On September 24, 2026, the Department of Justice lodged a proposed third consent decree modification with the United States District Court for the District of New Jersey in the lawsuit

entitled *United States v. The Sherwin-Williams Company*, Civil Action No. 1:19-cv-01907-ESK-EAP, an action brought under the Comprehensive Environmental Response, Compensation and Liability Act ("CERCLA"), 42 U.S.C. 9601 *et seq.*

The Court entered a consent decree ("Decree") in April 2019 in this action which resolved the United States' claim against The Sherwin-Williams Company ("Sherwin-Williams") for United States past costs relating to the Sherwin-Williams/Hilliards Creek Site, Route 561 Dump Site, and United States Avenue Burn Site (the "Sites"), located in Gibbsboro and Voorhees, New Jersey. The Decree also provides for Sherwin-Williams' performance of the soils and sediments operable unit remedy selected by the U.S. Environmental Protection Agency ("EPA") for the United States Avenue Burn Site.

The Decree also provides that, after EPA selects a remedy in a Record of Decision ("ROD") for an additional operable unit ("OU") at any of the Sites, the Decree may be modified to add Sherwin-Williams' performance of the

newly selected OU remedy to the work required under the Decree. In September 2025, EPA issued a ROD for an OU remedy that addresses groundwater at the Sherwin-Williams/Hilliards Creek Site ("SW/HC OU3 remedy"). This remedy provides for in-situ treatment of deep groundwater at the former manufacturing plant area at the SW/HC Site with chemical oxidation, chemical reduction, or sequestration and enhanced bioremediation. The proposed third modification lodged with the Court on September 24, 2026, provides for Sherwin-Williams' performance of the SW/HC OU3 remedy.

The publication of this notice opens a period for public comment on the proposed third modification. Comments should be addressed to the Assistant Attorney General, Energy and Natural Resources Division, and should refer to *United States v. The Sherwin-Williams Company*, Civil Action No. 1-19-cv-01907-ESK-EAP, D.J. Ref. No. 90-11-3-09023/2. All comments must be submitted no later than thirty (30) days after the publication date of this notice.