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**Heather Achbach,**

*Federal Register Liaison Officer, Drug Enforcement Administration.*

[FR Doc. 2023-28015 Filed 12-20-23; 8:45 am]

**BILLING CODE 4410-09-P**

**DEPARTMENT OF JUSTICE**

[OMB Number 1117-0003]

**Agency Information Collection Activities; Proposed eCollection Activities; Proposed eCollection Comments Requested; Extension of a Previously Approved Collection; ARCOS Transaction Reporting**

**AGENCY:** Drug Enforcement Administration, Department of Justice.  
**ACTION:** 60-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 60 days until February 20, 2024.

**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 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 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:* Extension of a previously approved collection.
2. *The Title of the Form/Collection:* ARCOS Transaction Reporting.
3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* DEA Form 333. 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: Private Sector—business or other for-profit. The obligation to respond is mandatory per 21 CFR 1304.
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 1,181 registrants participate in this information collection. The time per response is 0.50 minutes to complete the DEA-333 (paper) and 0.25 minutes to complete DEA-333 (online).
6. *An estimate of the total annual burden (in hours) associated with the collection:* DEA estimates that this collection takes 2,850 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	Total annual responses	Time per response (hours)	Total annual burden (hours)
DEA Form: 333 (online) .....	31	110	0.50	55
DEA Form: 333 (paper) .....	1,150	11,180	0.25	2,795
Unduplicated Totals .....	1,181	11,290	.....	2,850

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

Dated: December 18, 2023.

**Darwin Arceo,**

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

[FR Doc. 2023-28107 Filed 12-20-23; 8:45 am]

**BILLING CODE 4410-09-P**

**DEPARTMENT OF JUSTICE**

**Notice of Extension of Comment Period on Proposed Consent Decrees Under the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA), the Clean Water Act (CWA), and the Oil Pollution Act (OPA)**

On November 1, 2023, the Department of Justice lodged two proposed consent decrees with the United States District Court for the District of Oregon in the lawsuit entitled *United States of America et al. v. ACF Industries LLC, et al.*, Civil Action No. 3:23-cv-1603 (D. Or.). Notice of this settlement was published in the **Federal Register** at 88 FR 78063 (Nov. 14, 2023), which announced a 45-day comment period. Based on the date of that **Federal Register** notice, the comment period was scheduled to end on December 29, 2023.

On December 11, 2023, Plaintiffs in the above-captioned settlement received a request to extend the comment period by an additional forty-five (45) days. After considering this request, Plaintiffs have decided to extend the original comment period by an additional thirty (30) days. This extension provides a total comment period of seventy-five (75) days, through and including January 28, 2024.

Comments on the proposed Consent Decrees should be addressed to the Assistant Attorney General, Environment and Natural Resources Division, and should refer to *United States of America et al. v. ACF Industries LLC, et al.*, D.J. Ref. No. 90-11-2-06787/2.

Comments may be submitted either by email or by mail:

<i>To submit comments:</i>	<i>Send them to:</i>
By email .....	<i>pubcomment-ees.enrd@usdoj.gov.</i>

<i>To submit comments:</i>	<i>Send them to:</i>
By mail .....	Assistant Attorney General, U.S. DOJ—ENRD, P.O. Box 7611, Washington, DC 20044-7611.

During the public comment period, the Consent Decrees may be examined and downloaded at this Justice Department website: <https://www.justice.gov/enrd/consent-decree/us-et-al-v-acf-industries-llc-et-al>. Please note that this website contains the corrected version of the cash-out consent decree but not the version originally lodged with the court. The corrected version of the cash-out consent decree adds a legal entity for one of the settling defendants that inadvertently was omitted but does not change the scope of the operations covered by the consent decree or the amounts to be paid under the consent decree. Please refer to the corrected version of the cash-out consent decree when submitting comments. We will provide a paper copy of the Consent Decrees upon written request and payment of reproduction costs. Please mail your request and payment to: Consent Decree Library, U.S. DOJ—ENRD, P.O. Box 7611, Washington, DC 20044-7611.

Please enclose a check or money order for \$45.25 (without attachments) or \$631.25 (with attachments) (25 cents per page reproduction cost) payable to the United States Treasury.

**Kathryn C. Macdonald,**

*Assistant Section Chief, Environmental Enforcement Section, Environment and Natural Resources Division.*

[FR Doc. 2023-28017 Filed 12-20-23; 8:45 am]

**BILLING CODE 4410-15-P**

**DEPARTMENT OF JUSTICE**

**[OMB Number 1117-0021]**

**Agency Information Collection Activities; Proposed eCollection eComments Requested; Extension of a Previously Approved Collection; Dispensing Records of Individual Practitioners**

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

**ACTION:** 60-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 60 days until February 20, 2024.

**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 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 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. 827(c), practitioners who regularly dispense or administer controlled substances to patients and charge them for the substances and those practitioners who administer controlled substances in the course of maintenance or detoxification treatment shall keep records of such activities, and accordingly must comply with the regulations on recordkeeping.

**Overview of This Information Collection**

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