DEPARTMENT OF JUSTICE

Bureau of Alcohol, Tobacco, Firearms and Explosives

[OMB Number 1140–0087]

Agency Information Collection Activities; Proposed eCollection of eComments Requested; Extension Without Change of a Currently Approved Collection; eForm Access Request/User Registration

AGENCY: Bureau of Alcohol, Tobacco, Firearms and Explosives, Department of Justice.

ACTION: 30-Day notice.

SUMMARY: The Bureau of Alcohol, Tobacco, Firearms and Explosives (ATF), Department of Justice (DOJ) will submit 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 an additional 30 days until September 16, 2021.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *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.

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;

—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.

Overview of This Information Collection

(1) *Type of Information Collection:* Extension without change of a currently approved collection.

(2) *The Title of the Form/Collection:* eForm Access Request/User Registration.

(3) *The agency form number, if any, and the applicable component of the Department sponsoring the collection:*

Form number: None.

Component: Bureau of Alcohol, Tobacco, Firearms and Explosives, U.S. Department of Justice.

(4) *Affected public who will be asked or required to respond, as well as a brief abstract:*

Primary: Business or other for profit.

Other: None.

Abstract: Members of the public will use the eForm Access Request/User Registration to create a username and password for access to the Bureau of Alcohol, Tobacco, Firearms, and Explosives' (ATF's) eForms platform, which is an electronic application filing system.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* An estimated 76,000 respondents will complete this registration form annually, and it will take each respondent approximately 2.24 minutes to complete their responses.

(6) *An estimate of the total public burden (in hours) associated with the collection:* The estimated annual public burden associated with this collection is 2,387 hours, which is equal to 76,000 (# of respondents) * .037333333 (2.24 minutes).

If additional information is required contact: Melody Braswell, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Two Constitution Square, 145 N Street NE, Mail Stop 3E.405A, Washington, DC 20530.

Dated: August 12, 2021.

Melody Braswell,

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

[FR Doc. 2021–17611 Filed 8–16–21; 8:45 am]

BILLING CODE 4410–FY–P

DEPARTMENT OF JUSTICE

[OMB Number 1117–0049]

Agency Information Collection Activities: Proposed eCollection, eComments Requested; Reinstatement of a Discontinued Collection: Recordkeeping for Electronic Prescriptions for Controlled Substances

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 October 18, 2021.

FOR FURTHER INFORMATION CONTACT: If you have comments 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, Diversion Control Division, Drug Enforcement Administration; Mailing Address: 8701 Morrissette Drive, Springfield, Virginia 22152; Telephone: (571) 362–3261.

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;

—Evaluate whether and if so how the quality, utility, and clarity of the information proposed 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 forms of information technology, *e.g.,* permitting electronic submission of responses.

Overview of This Information Collection

1. *Type of Information Collection:* Reinstatement of a discontinued collection.

2. *Title of the Form/Collection:* Recordkeeping for Electronic Prescriptions for Controlled Substance.

3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* There is no form number. 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): Business or other for-profit.

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

Abstract: DEA is requiring that each registered practitioner apply to an approved credential service provider approved to obtain identity proofing and a credential. Hospitals and other institutional practitioners may conduct this process in-house as part of their credentialing. For practitioners currently working at or affiliated with a registered hospital or clinic, the hospital/clinic have to check a government-issued photographic identification. This may be done when the hospital/clinic issues credentials to new hires or newly affiliated physicians. For individual practitioners, two people need to enter logical access control data to grant permissions for practitioners authorized to approve and sign controlled substance prescriptions using the electronic prescription application. For institutional practitioners, logical access control data is entered by two people from an entity within the hospital/clinic that is separate from the entity that conduct identity proofing in-house. Similarly, pharmacies have to set logical access controls in the pharmacy application so that only authorized employees have permission to annotate or alter prescription records. Finally, if the electronic prescription or pharmacy application generates an incident report, practitioners, hospitals/clinics, and pharmacies have to review the incident report to determine if the event identified by the application represents a security incident.

5. *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* The below table presents information regarding the number of respondents, hour burden per responses and associated burden hours.

	Number of respondents	Hour burden per response	Burden hours
Practitioners	78,164	0.67	52,370
MLP	49,067	0.67	32,875
Hospital/Clinics	1,482	2.13	3,157
Pharmacies	3,984	0.33	1,315
Total	132,697		89,717

6. *An estimate of the total public burden (in hours) associated with the proposed collection:* DEA estimates that this collection takes 89,717 annual burden hours.

If additional information is required please contact: Melody Braswell, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Two Constitution Square, 145 N Street NE, Suite 3E.405B, Washington, DC 20530.

Dated: August 11, 2021.

Melody Braswell,

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

[FR Doc. 2021–17523 Filed 8–16–21; 8:45 am]

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DEPARTMENT OF LABOR

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Respiratory Protection Standard

ACTION: Notice of availability; request for comments.

SUMMARY: The Department of Labor (DOL) is submitting this Occupational Safety and Health Administration (OSHA)-sponsored information collection request (ICR) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995 (PRA). Public comments on the ICR are invited.

DATES: The OMB will consider all written comments that agency receives on or before September 16, 2021.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *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.

Comments are invited on: (1) Whether the collection of information is necessary for the proper performance of the functions of the Department, including whether the information will have practical utility; (2) if the information will be processed and used in a timely manner; (3) the accuracy of the agency's estimates of the burden and cost of the collection of information, including the validity of the methodology and assumptions used; (4) ways to enhance the quality, utility and clarity of the information collection; and (5) ways to minimize the burden of the collection of information on those who are to respond, including the use of automated collection techniques or other forms of information technology.

FOR FURTHER INFORMATION CONTACT: Crystal Rennie by telephone at 202–693–0456 or by email at *DOL_PRA_PUBLIC@dol.gov.*

SUPPLEMENTARY INFORMATION: This standard requires employers to develop a written respiratory protection program, provide medical surveillance, fit test employees, obtain certificates of analysis on cylinders, change sorbent beds and filters, to inspect emergency-use respirators, mark emergency-use respirator storage compartments, and maintain accurate employee records for fit testing and medical surveillance. For additional substantive information about this ICR, see the related notice published in the **Federal Register** on April 9, 2021 (86 FR 18557).

This information collection is subject to the PRA. A Federal agency generally cannot conduct or sponsor a collection of information, and the public is generally not required to respond to an information collection, unless the OMB approves it and displays a currently