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 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:** Revision of a currently approved collection.

2. **Title of the Form/Collection:** Application for Registration under Domestic Chemical Diversion Control Act of 1993; Renewal Application for Registration under Domestic Chemical Diversion Control Act of 1993.

3. **The agency form number, if any, and the applicable component of the Department sponsoring the collection:** DEA Forms: 510, 510A. 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): None. Abstract: The DEA implements the Controlled Substances Act (CSA) which requires that every person who manufactures or distributes a list I chemical shall annually obtain a registration for that purpose. The DEA will be revising the proposed information collection instruments concerning the liability questions on the Application for Registration under Domestic Chemical Diversion Control Act of 1993; and Renewal Application for Registration under Domestic Chemical Diversion Control Act of 1993. Over the years, many applicants have answered some of the liability questions incorrectly. These changes will avoid confusion to the applicant by separating compound questions into multiple parts that will require the applicant to answer them individually.

5. **An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:**

| DEA–510 (paper) | 56 | 0.2 hours (12 minutes) | 1.20 |
| DEA–510 (electronic) | 228 | 0.17 hours (8 minutes) | 11.73 |
| DEA–510A (paper) | 28 | 0.2 hours (10 minutes) | 4.67 |
| DEA–510A (electronic) | 874 | 0.07 hours (4 minutes) | 58.27 |
| **Total** | **996** | | **76.87** |

6. **An estimate of the total public burden (in hours) associated with the proposed collection:** The DEA estimates that this collection takes 76.87 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.


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

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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; 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 Department of Justice, Drug Enforcement Administration (DEA), will be submitting the following information collection request to the Office of Management and Budget 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 12, 2019.

**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 Kathy L. Federico, Diversion Control Division, Drug Enforcement Administration; Mailing Address: 8701 Morrissette Drive, Springfield, Virginia 22152; Telephone: (202) 598–6812.

**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 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 of a currently approved collection.

2. Title of the Form/Collection: Application for Individual Manufacturing Quota for a Basic Class of 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: DEA Form 189. 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): None.

ABSTRACT: Pursuant to 21 U.S.C. 826(c) and 21 CFR 1303.22 and 1315.22, any person who is registered to manufacture any basic class of controlled substances listed in Schedule I or II, or the List I chemicals ephedrine, pseudoephedrine, or phenylpropanolamine, and who desires to manufacture a quantity of such class or such List I chemical, must apply on DEA Form 189 for a manufacturing quota for such quantity of such class or List I chemical.

5. An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond: The DEA estimates 33 respondents complete 859 DEA Form 189 applications annually, and that each form takes 0.5 hours to complete.

6. An estimate of the total public burden (in hours) associated with the proposed collection: The DEA estimates this collection takes a total of 430 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.


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

DEPARTMENT OF LABOR
Employment and Training Administration

Agency Information Collection Activities; Comment Request; Statement of Expenditures and Financial Adjustments of Federal Funds for Unemployment Compensation for Federal Employees and Ex-Servicemembers Report

ACTION: Notice.

SUMMARY: The Department of Labor’s (DOL’s) Employment and Training Administration (ETA) is soliciting comments concerning a proposed extension for the authority to conduct the information request (ICR) titled, “Statement of Expenditures and Financial Adjustments of Federal Funds for Unemployment Compensation for Federal Employees and Ex-Servicemembers Report.” This comment request is part of continuing Departmental efforts to reduce paperwork and respondent burden in accordance with the Paperwork Reduction Act of 1995 (PRA).

DATES: Consideration will be given to all written comments received by February 12, 2019.

ADDRESSES: A copy of this ICR with applicable supporting documentation; including a description of the likely respondents, proposed frequency of response, and estimated total burden may be obtained free by contacting Cindy Le by telephone at (202) 693–2829, TTY 1–877–889–5627 (these are not toll-free numbers), or by email at le.cindy@dol.gov.

Submit written comments about or requests for a copy of this ICR by mail or courier to the U.S. Department of Labor, Employment and Training Administration, Office of Unemployment Insurance, Room S–4524, 200 Constitution Avenue NW, Washington, DC 20210, by email to le.cindy@dol.gov, or by Fax at (202) 693–3975.

SUPPLEMENTARY INFORMATION: As part of continuing efforts to reduce paperwork and respondent burden, DOL conducts a pre-clearance consultation program to provide the general public and Federal agencies an opportunity to comment on proposed and/or continuing collections of information before submitting them to the Office of Management and Budget (OMB) for final approval. This program helps to ensure requested data is provided in the desired format, reporting burden (time and financial resources) is minimized, collection instruments are clearly understood, and the impact of collection requirements can be properly assessed.

Public Law 97–362, Miscellaneous Revenue Act of 1982, amended the Unemployment Compensation for Ex-Servicemembers (UCX) law (5 U.S.C. 8509), and Public Law 96–499, Omnibus Budget Reconciliation Act, amended the Unemployment Compensation for Federal Employees (UCFE) law (5 U.S.C. 8501, et seq.), requiring each Federal employing agency to pay the costs of regular and extended UCFE/UCX benefits paid to its employees by the State Workforce Agencies (SWAs). The ETA 191 report submitted quarterly by each SWA shows the amounts of benefits that should be charged to each Federal employing agency. The Office of Unemployment Insurance uses this information to aggregate the SWA quarterly charges and submit one official bill to each Federal agency being charged. Federal agencies then reimburse the Federal Employees Compensation Account maintained by the U.S. Treasury. This collection is authorized by the Social Security Act, Section 303(a)(6).

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 it is approved by OMB under the PRA and displays a currently valid OMB Control Number. In addition, notwithstanding any other provisions of law, no person shall generally be subject to penalty for failing to comply with a collection of information that does not display a valid Control Number. See 5 CFR 1320.5(a) and 1320.6.

Interested parties are encouraged to provide comments to the contact shown in the ADDRESSES section. Comments must be written to receive consideration, and they will be summarized and included in the request for OMB approval of the final ICR. In order to help ensure appropriate consideration, comments should mention OMB control number 1205–0162.

Submitted comments will also be a matter of public record for this ICR and posted on the internet, without redaction. DOL encourages commenters not to include personally identifiable information, confidential business data, or other sensitive statements/information in any comments.

DOL is particularly interested in comments that:
• Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including