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.

Dated: March 2, 2022.

Melody Braswell,

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

[FR Doc. 2022–04786 Filed 3–7–22; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

[OMB Number 1117–0008]

Agency Information Collection Activities; Proposed eCollection, eComments Requested; Extension Without Change of a Previously Approved Collection; Application for Procurement Quota for Controlled Substance and for Ephedrine, Pseudoephedrine, and Phenylpropanolamine; DEA Form 250

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. This information collection is also associated with the proposed rulemaking ''Management of Quotas for Controlled Substances and List I Chemicals,'' published in the **Federal Register**. It is likely that the final rule will not be published before this information collection expires on June 30, 2022. If the final rule does publish prior to the expiration, it will be published as the 30-Day Notice.

DATES: Comments are encouraged and will be accepted for 60 days until May 9, 2022.

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 (DPW), Diversion Control Division, Drug Enforcement Administration; Mailing Address: 8701 Morrissette Drive, Springfield, Virginia 22152; Telephone: (571) 776–2265.

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 Procurement Quota for 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 250. 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 and 21 CFR 1303.12(b) and 1315.32, any person who desires to use, during the next calendar year, any basic class of controlled substances listed in schedules I or II, or the List I chemicals ephedrine, pseudoephedrine, or phenylpropanolamine for purposes of manufacturing must apply on DEA Form 250 for a procurement quota for 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 344 respondents complete 3,066 DEA Form 250 applications annually, and that each form requires 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 1,533 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: March 2, 2022.

Melody Braswell,

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

[FR Doc. 2022–04785 Filed 3–7–22; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

[OMB Number 1117–0047]

Agency Information Collection Activities; Proposed eCollection, eComments Requested; Extension Without Change of a Previously Approved Collection; Application for Import Quota for Ephedrine, Pseudoephedrine, and Phenylpropanolamine; DEA Form 488

AGENCY: Drug Enforcement Administration, Department of Justice.

ACTION: 60-Day notice.

SUMMARY: The Drug Enforcement Administration (DEA), Department of Justice, 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. This information collection is also associated with the proposed rulemaking ''Management of Quotas for Controlled Substances and List I Chemicals,'' published in the **Federal Register**. It is likely that the final rule will not be published before this information collection expires on May 31, 2022. If the final rule does publish prior to the expiration, it will be published as the 30-Day Notice.

DATES: Comments are encouraged and will be accepted for 60 days until May 9, 2022.

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 (DPW), Diversion Control Division, Drug Enforcement Administration; Mailing Address: 8701 Morrissette Drive, Springfield, Virginia 22152; Telephone: (571) 776–2265.

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 Import Quota for Ephedrine, Pseudoephedrine, and Phenylpropanolamine.

3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* DEA Form 488. 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: Pursuant to 21 U.S.C. 952 and 21 CFR 1315.34, any person who desires to import the List I chemicals Ephedrine, Pseudoephedrine, or Phenylpropanolamine during the next calendar year must apply on DEA Form 488 for an import quota for each such 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 49 respondents complete 126 DEA Form 488 applications annually, and that each form takes 0.5 hours to complete. Respondents complete a separate DEA Form 488 for each List I chemical for which quota is sought.

6. *An estimate of the total public burden (in hours) associated with the proposed collection:* The DEA estimates this collection takes a total of 63 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: March 2, 2022.

Melody Braswell,

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

[FR Doc. 2022–04787 Filed 3–7–22; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Notice of Lodging Proposed Consent Decree

In accordance with Departmental Policy, 28 CFR 50.7, notice is hereby given that a proposed Consent Decree in *United States* v. *Polo Development, Inc., et al.,* Civil Action No. 4:20–cv–2400–JRA, was lodged with the United States District Court for the Northern District of Ohio on March 1, 2022.

This proposed Consent Decree concerns an amended complaint filed by the United States against Defendants Polo Development, Inc., AIM Georgia, LLC, Joseph Zdrilich, Donna Zdrilich, and Carbon Hills, LLC, pursuant to Section 309(b) of the Clean Water Act, 33 U.S.C. 1319(b), to obtain injunctive relief from and impose civil penalties against the Defendants for violating Section 301(a) of the Clean Water Act, 33 U.S.C. 1311(a), by discharging pollutants without a permit into waters of the United States. The proposed Consent Decree resolves these claims by requiring the Defendants to restore impacted areas, record a conservation easement, and pay a civil penalty.

The Department of Justice will accept written comments relating to this proposed Consent Decree for thirty (30) days from the date of publication of this Notice. Please address comments to Patrick R. Jacobi, United States Department of Justice, Environment and Natural Resources Division, Environmental Defense Section, Denver Place Building, 999 18th Street, Suite 370—South Terrace, Denver, CO 80202, *pubcomment_eds.enrd@usdoj.gov,* and refer to *United States* v. *Polo Development, Inc., et al.,* DJ #'s 90–5–1–1–21099, 90–5–1–1–22034.

Subject to public health protocols, the proposed Consent Decree may be