958, to the DEA Administrator, 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been delegated to the Assistant Administrator of the DEA Diversion Control Division (''Assistant Administrator'') pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

Therefore, in accordance with 21 U.S.C. 958(i) and 21 CFR 1301.34(a), this is notice that on June 11, 2018, Arizona Department of Corrections, 1305 E Butte Avenue, ASPC-Florence, Florence, Arizona 85132–9221, re-applied to be registered as an importer of Pentobarbital (2270), a basic class of the controlled substance listed in schedule II.

The facility intends to import the above-listed controlled substance for legitimate use. This particular controlled substance is not available for the intended legitimate use within the current domestic supply of the United States.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture this basic class of controlled substance may file comments or objections to the issuance of the proposed registration or to the authorization of this importation, and may, at the same time, file a written request for a hearing. Any such comments, objections, or hearing requests should be addressed as described above.

Dated: December 4, 2018.

John J. Martin,

Assistant Administrator.

[FR Doc. 2018–27131 Filed 12–13–18; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: Usona Institute

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before January 14, 2019. Such persons may also file a written request for a hearing on the application on or before January 14, 2019.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division (''Assistant Administrator'') pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on October 31, 2018, Usona Institute, 2800 Woods Hollow Road, Madison, Wisconsin 53711 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
5-Methoxy-N-N-dimethyltryptamine	7431	I
Dimethyltryptamine	7435	I

The institute plans to import the listed controlled substances for potential formulation development for substances to be used in institute-sponsored research.

Dated: December 4, 2018.

John J. Martin,

Assistant Administrator.

[FR Doc. 2018–27133 Filed 12–13–18; 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.

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 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: December 10, 2018.

Melody Braswell,

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

[FR Doc. 2018–27059 Filed 12–13–18; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

[OMB Number 1117–0029]

Agency Information Collection Activities; Proposed eCollection, eComments Requested; Extension Without Change of a Previously Approved Collection; Annual Reporting Requirement for Manufacturers of Listed Chemicals

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:* Annual Reporting Requirement for Manufacturers of Listed Chemicals.

3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* Form number: N/A. 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. 830(b)(2) and 21 CFR 1310.05(d), manufacturers of listed chemicals must file annual reports of manufacturing, inventory, and use data for the listed chemicals they manufacture. These reports allow the DEA to monitor the volume and availability of domestically manufactured listed chemicals, which may be subject to diversion for the illicit production of controlled substances.

5. *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* Each respondent for this information collection completes one response per year. The DEA estimates there are 50 respondents, and that each response takes 0.25 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 12.5 annual burden hours.