

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.

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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 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement

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

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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 Morrisette 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