

service list), and a certificate of service must be timely filed. The Secretary will not accept a document for filing without a certificate of service.

Authority: These investigations are being conducted under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to § 207.21 of the Commission's rules.

By order of the Commission.

Issued: June 9, 2022.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2022-12953 Filed 6-17-22; 8:45 am]

BILLING CODE 7020-02-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1026]

Importer of Controlled Substances

Application: Alcami Carolinas Corporation

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Alcami Carolinas Corporation has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to Supplementary Information listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before July 21, 2022. Such persons may also file a written request for a hearing on the application on or before July 21, 2022.

ADDRESSES: The Drug Enforcement Administration (DEA) requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to <https://www.regulations.gov> and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on <https://www.regulations.gov>. If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment. All requests for a hearing must 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. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on May 10, 2022, Alcami Carolinas Corporation, 1726 North 23rd Street, Attn: DEA Compliance Office, Wilmington, North Carolina 28405-1827, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Psilocybin	7437	I
Psilocyn	7438	I
Pentobarbital	2270	II
Thebaine	9333	II

The company plans to import the listed controlled substances in bulk for the manufacturing of capsules/tablets for Phase II clinical trials. The company plans to import derivatives of Thebaine that have been determined by DEA to be captured under drug code (9333) Thebaine. No other activity for these drug codes are authorized for this registration.

Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

Kristi O'Malley,

Assistant Administrator.

[FR Doc. 2022-13246 Filed 6-17-22; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1028]

Importer of Controlled Substances

Application: Arizona Department of Corrections

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Arizona Department of Corrections has applied to be registered as an importer of basic class(es) of

controlled substance(s). Refer to Supplementary Information listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before July 21, 2022. Such persons may also file a written request for a hearing on the application on or before July 21, 2022.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to <https://www.regulations.gov> and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on <https://www.regulations.gov>. If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment. All requests for a hearing must 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. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on May 13, 2022, Arizona Department of Corrections, 1305 East Butte Avenue, ASPC-Florence, Florence, Arizona 85132-9221, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Pentobarbital	2270	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. No other activity for these drug codes are authorized for this registration.

Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

Kristi O'Malley,
Assistant Administrator.
 [FR Doc. 2022-13248 Filed 6-17-22; 8:45 am]
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DEPARTMENT OF JUSTICE

Notice of Lodging of Proposed Stipulation and Order of Settlement Under the Federal Insecticide, Fungicide, and Rodenticide Act

On June 13, 2022, the Department of Justice lodged a proposed Stipulation and Order of Settlement ("settlement") with the District Court of the Southern District of New York in a lawsuit entitled *Tzumi Innovations, LLC v. Regan, et al.*, Civil Action No. 21-122.

In a counterclaim in this action, the United States seeks, as provided under the Federal Insecticide, Fungicide, and Rodenticide Act ("FIFRA"), a civil penalty and injunctive relief from Tzumi Innovations, LLC ("Tzumi") in connection with Tzumi's distribution and sale of unregistered pesticides pursuant to FIFRA § 12(a)(1)(A), 7 U.S.C. 136j(a)(1)(A). The proposed settlement resolves the United States' counterclaim, requires Tzumi to pay \$1.5 million, and imposes injunctive relief. It also resolves Tzumi's claims against the Environmental Protection Agency in the litigation.

The publication of this notice opens the public comment on the proposed settlement. Comments should be addressed to the Assistant Attorney General, Environment and Natural Resources Division, and should refer to *Tzumi Innovations, LLC v. Regan, D.J. # 1-12610*. All comments must be submitted no later than 30 days after the publication date of this notice. Comments may be submitted either by email or by mail:

<i>To submit comments:</i>	<i>Send them to:</i>
By email	<i>pubcomment-ees.enrd@usdoj.gov.</i>
By mail	Assistant Attorney General, U.S. DOJ—ENRD, P.O. Box 7611, Washington, DC 20044-7611.

During the public comment period, the settlement may be examined and

downloaded at this Justice Department website: http://www.usdoj.gov/enrd/Consent_Decrees.html. We will provide a paper copy of the settlement upon written request and payment of reproduction costs. Please email your request and payment to: Consent Decree Library, U.S. DOJ—ENRD, P.O. Box 7611, Washington, DC 20044-7611.

Please enclose a check or money order for \$6.50 (25 cents per page reproduction cost) payable to the United States Treasury.

Henry S. Friedman,
Assistant Section Chief, Environmental Enforcement Section, Environment and Natural Resources Division.
 [FR Doc. 2022-13167 Filed 6-17-22; 8:45 am]
BILLING CODE 4410-15-P

DEPARTMENT OF LABOR

Agency Information Collection Activities; OSHA Strategic Partnership Program (OSSP) for Worker Safety and Health

ACTION: Notice of availability; request for comments.

SUMMARY: The Department of Labor (DOL) is submitting this Occupational Safety & 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 the agency receives on or before July 21, 2022.

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) 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; (3) ways to enhance the quality, utility and clarity of the information collection; and (4) 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: Nicole Bouchet by telephone at 202-693-0213, or by email at DOL_PRA_PUBLIC@dol.gov.

SUPPLEMENTARY INFORMATION: The Occupational Safety and Health Act of 1970 (OSH Act) (29 U.S.C. 651 *et seq.*) authorizes information collection by employers as necessary or appropriate for enforcement of the OSH Act or for developing information regarding the causes and prevention of occupational injuries, illnesses, and accidents (29 U.S.C. 657). The OSH Act also requires that OSHA obtain such information with a minimum burden upon employers, especially those operating small businesses, and to reduce to the maximum extent feasible unnecessary duplication of efforts in obtaining said information (29 U.S.C. 657). The OSPP allows OSHA to enter into an extended, voluntary, cooperative relationship with groups of employers, employees, and representatives (sometimes including other stakeholders, and sometimes involving only one employer) to encourage, assist, and recognize their efforts to eliminate serious hazards and to achieve a high level of worker safety and health that goes beyond what historically has been achieved from traditional enforcement methods. Each OSHA Strategic Partnership (OSP) determines what information will be needed, determining the best collection method, and clarifying how the information will be used. At a minimum, each OSP must identify baseline injury and illness data corresponding to all summary line items on the OSHA 300 logs and must track changes at either the worksite level or participant-aggregate level. An OSP may also include other measures of success, such as training activity, self-inspections, and/or workers' compensation data. In this regard, the information collection requirements for the OSPP are used by the agency to gauge the effectiveness of programs, identify needed improvements, and ensure that resources are being used effectively and appropriately. For additional substantive information about this ICR, see the related notice published in the **Federal Register** on March 7, 2022 (87 FR 12735).

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