

Company	FR docket	Published
Meridian Medical Technologies	84 FR 7129	March 1, 2019.
Organic Standards Solutions International, LLC	84 FR 13958	April 8, 2019.
SpecGx LLC	84 FR 13954	April 8, 2019.

The DEA has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of the listed registrants to import the applicable basic classes of schedule I and II controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated each of the company's maintenance of effective controls against diversion by inspecting and testing each company's physical security systems, verifying each company's compliance with state and local laws, and reviewing each company's background and history.

Therefore, pursuant to 21 U.S.C. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the DEA has granted a registration as an importer for schedule I and schedule II controlled substances to the above listed companies.

Dated: May 7, 2019.

John J. Martin,

Assistant Administrator.

[FR Doc. 2019-10669 Filed 5-21-19; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

**Importer of Controlled Substances
Application: United States
Pharmacoepial Convention**

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 June 21, 2019. Such persons may also file a written request for a hearing on the application on or before June 21, 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 a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All request for a 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 March 8, 2019, United States Pharmacoepial, 12601 Twinbrook Parkway, Rockville, Maryland 20852-1717 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Methamphetamine	1105	II
Cathinone	1235	I
Phenmetrazine	1631	II
Methylphenidate	1724	II
Amobarbital	2125	II
Pentobarbital	2270	II
Secobarbital	2315	II
Glutethimide	2550	II
Methaqualone	2565	I
Lysergic acid diethylamide	7315	I
4-Methyl-2,5-dimethoxyamphetamine	7395	I
3,4-Methylenedioxyamphetamine	7400	I
4-Methoxyamphetamine	7411	I
Phencyclidine	7471	II
4-Anilino-N-phenethyl-4-piperidine (ANPP)	8333	II
Phenylacetone	8501	II
Alphaprodine	9010	II
Anileridine	9020	II
Cocaine	9041	II
Codeine-N-oxide	9053	I
Dihydrocodeine	9120	II
Difenoxin	9168	I
Diphenoxylate	9170	II
Heroin	9200	I
Levomethorphan	9210	II
Levorphanol	9220	II
Meperidine	9230	II
Dextropropoxyphene, bulk (non-dosage forms)	9273	II

Controlled substance	Drug code	Schedule
Morphine-N-oxide	9307	I
Thebaine	9333	II
Norlevorphanol	9634	II
Oxymorphone	9652	II
Noroxymorphone	9668	II
Alfentanil	9737	II
Sufentanil	9740	II

The company plans to import the bulk control substances for distribution of analytical reference standards to its customers for analytical testing of raw materials.

Dated: May 7, 2019.

John J. Martin,

Assistant Administrator.

[FR Doc. 2019-10668 Filed 5-21-19; 8:45 am]

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DEPARTMENT OF JUSTICE

Foreign Claims Settlement Commission

[F.C.S.C. Meeting and Hearing Notice No. 04-19]

Sunshine Act Meeting

The Foreign Claims Settlement Commission, pursuant to its regulations (45 CFR part 503.25) and the Government in the Sunshine Act (5 U.S.C. 552b), hereby gives notice in regard to the scheduling of open meetings as follows:

TIME AND DATE: Thursday, May 30, 2019, at 10:00 a.m.

PLACE: All meetings are held at the Foreign Claims Settlement Commission, 601 D Street NW, Suite 10300, Washington, DC.

STATUS: Open.

MATTERS TO BE CONSIDERED: 10:00 a.m.—Oral hearings on Objection to Commission’s Proposed Decisions in Claim Nos. IRQ-II-346 and IRQ-II-365.

11:30 a.m.—Issuance of Proposed Decisions under the Guam World War II Loyalty Recognition Act, Title XVII, Public Law 114-328.

CONTACT PERSON FOR MORE INFORMATION: Requests for information, or advance notices of intention to observe an open meeting, may be directed to: Patricia M. Hall, Foreign Claims Settlement Commission, 601 D Street NW, Suite 10300, Washington, DC 20579. Telephone: (202) 616-6975.

Brian Simkin,
Chief Counsel.

[FR Doc. 2019-10735 Filed 5-20-19; 11:15 am]

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DEPARTMENT OF JUSTICE

Notice of Lodging of Proposed Settlement Agreement Under the Clean Air Act

On May 16, 2019, the Department of Justice lodged a proposed Stipulation of Settlement and Order (“Agreement”) with the United States District Court for the District of Massachusetts in the lawsuit entitled *United States v. Kayem Foods, Inc.*, Civil Action No. 1:19-cv-11126.

In this action, the United States filed a complaint alleging that Kayem Foods, Inc. (“Kayem”) violated Section 112(r)(7) of the Clean Air Act, 42 U.S.C. 7412(r)(7), at Kayem’s food processing facility located in Chelsea, Massachusetts. Section 112(r)(7) of the CAA, 42 U.S.C. 7412(r)(7), provides that the Administrator of the EPA is authorized to promulgate regulations requiring owners or operators of a stationary source at which a regulated substance is present in more than a threshold amount to, among other things, prepare and implement a risk management plan to detect and prevent or minimize accidental releases of regulated substances from the stationary source, and to provide a prompt emergency response to any such releases in order to protect human health and the environment. EPA has promulgated regulations to implement Section 112(r)(7), codified at 40 CFR part 68 (“Part 68 Regulations”). The Complaint alleges that Kayem violated the Part 68 Regulations in connection with the operation of its ammonia refrigeration system at its Chelsea facility and seeks the payment of civil penalties.

The proposed Agreement resolves Kayem’s civil liability to the United States for the alleged violations in the Complaint. Pursuant to the proposed Agreement, Kayem will pay a penalty of \$138,281. Injunctive relief is not required.

The publication of this notice opens a period for public comment on the Agreement. Comments should be addressed to the Assistant Attorney General, Environment and Natural Resources Division, and should refer to

United States v. Kayem Foods, Inc., No. 1:19-cv-11126 (D. Mass.) D.J. Ref. No. 90-5-2-1-11490. All comments must be submitted no later than thirty (30) days after the publication date of this notice. Comments may be submitted either by email or by mail:

To submit comments:	Send them to:
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 Agreement may be examined and downloaded at this Justice Department website: <https://www.justice.gov/enrd/consent-decrees>. We will provide a paper copy of the Agreement upon written request and payment of reproduction costs. Please mail 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 \$2.75 (25 cents per page reproduction cost) payable to the United States Treasury.

Robert Maher,

Assistant Section Chief, Environmental Enforcement Section, Environment and Natural Resources Division.

[FR Doc. 2019-10673 Filed 5-21-19; 8:45 am]

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DEPARTMENT OF LABOR

Employment and Training Administration

Agency Information Collection Activities; Comment Request; State Training Provider Eligibility Collection

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 collection request (ICR) titled, “State Training Provider