suspension agreements took effect on March 17, 2023. The antidumping duty suspension agreement is based upon an agreement between Commerce and producers/exporters which account for substantially all imports of white grape juice concentrate from Argentina, in which each signatory producer/exporter has agreed to revise its prices to eliminate completely the injurious effects of exports of WGJC to the United States. The countervailing duty suspension agreement is based upon an agreement between Commerce and the Government of Argentina (''GOA''), wherein the GOA has agreed not to provide any new or additional export or import substitution subsidies on the subject merchandise and has agreed to restrict the volume of direct or indirect exports to the United States of WGJC from all Argentine producers/exporters in order to eliminate completely the injurious effects of exports of this merchandise to the United States. Accordingly, the U.S. International Trade Commission gives notice of the suspension of its antidumping and countervailing duty investigations involving imports of WGJC from Argentina, provided for in subheading 2009.69.00 of the Harmonized Tariff Schedule of the United States.

DATES: March 24, 2023.

FOR FURTHER INFORMATION CONTACT: Ahdia Bavari (202–205–3191), Office of Investigations, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436. Hearing-impaired individuals are advised that information on this matter can be obtained by contacting the Commission's TDD terminal on 202–205–1810. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at 202–205–2000. General information concerning the Commission may also be obtained by accessing its internet server (*https://www.usitc.gov*). The public record for these investigations may be viewed on the Commission's electronic docket (EDIS) at *https://edis.usitc.gov.*

Authority: These investigations are being suspended under authority of title VII of the Tariff Act of 1930 and pursuant to section 207.40(b) of the Commission's Rules of Practice and Procedure (19 CFR 207.40(b)). This notice is published pursuant to section 201.10 of the Commission's rules (19 CFR 201.10).

By order of the Commission.

Issued: March 27, 2023.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2023–06669 Filed 3–30–23; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[USITC SE–23–018]

Sunshine Act Meetings

AGENCY HOLDING THE MEETING: United States International Trade Commission.

TIME AND DATE: April 7, 2023 at 11:00 a.m.

PLACE: Room 101, 500 E Street SW, Washington, DC 20436, Telephone: (202) 205–2000.

STATUS: Open to the public.

MATTERS TO BE CONSIDERED:

1. *Agendas for future meetings:* none.
2. Minutes.
3. Ratification List.
4. Commission vote on Inv. Nos. 701–TA–552 and 731–TA–1308 (Review) (Pneumatic Off-the-Road (OTR) Tires from India). The Commission currently is scheduled to complete and file its determinations and views of the Commission on April 27, 2023.
5. *Outstanding action jackets:* none.

CONTACT PERSON FOR MORE INFORMATION: Sharon Bellamy, Acting Supervisory Hearings and Information Officer, 202–205–2000.

The Commission is holding the meeting under the Government in the Sunshine Act, 5 U.S.C. 552(b). In accordance with Commission policy, subject matter listed above, not disposed of at the scheduled meeting, may be carried over to the agenda of the following meeting.

By order of the Commission.

Issued: March 29, 2023.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2023–06864 Filed 3–29–23; 4:15 pm]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1174]

Bulk Manufacturer of Controlled Substances Application: Sterling Pharma USA LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Sterling Pharma USA LLC. has applied to be registered as a bulk manufacturer 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 May 30, 2023. Such persons may also file a written request for a hearing on the application on or before May 30, 2023.

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.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on March 3, 2023, Sterling Pharma USA LLC., 1001 Sheldon Drive, Suite 101, Cary, North Carolina 27513–2078, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Dimethyltryptamine	7435	I

The company plans to manufacture the above-listed controlled substance(s) to support clinical trials. No other activities for this drug code is authorized for this registration.

Matthew Strait,

Deputy Assistant Administrator.

[FR Doc. 2023–06698 Filed 3–30–23; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1172]

Bulk Manufacturer of Controlled Substances Application: Purisys, LLC

AGENCY: Drug Enforcement Administration, Justice.