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 205.18-1, with the GOA has agreed to revise its prices to eliminate completely the injurious effects of exports of WGJC to the United States International Trade Commission.

MATTERS TO BE CONSIDERED:
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 7, 2023 at 11:00 a.m.
5. Outstanding action jackets: none.

CONTACT PERSON FOR MORE INFORMATION:
Sharon Bellamy. Acting Deputy Assistant Administrator.

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.

DEPARTMENT OF JUSTICE Drug Enforcement Administration
[Docket No. DEA–1172]
Bulk Manufacturer of Controlled Substances Application: Pharmaco USA LLC
AGENCY: Drug Enforcement Administration, Justice.
ACTION: Notice of application.
SUMMARY: Pharmaco 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, Pharmaco 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):

<table>
<thead>
<tr>
<th>Controlled substance</th>
<th>Drug code</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dimethyltryptamine</td>
<td>7435 I</td>
<td>1</td>
</tr>
</tbody>
</table>

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.