Commission, 500 E Street SW, Washington, DC 20436. Hearing-impaired persons can obtain information on this matter 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.

SUPPLEMENTARY INFORMATION: Effective February 24, 2021, the Commission established a general schedule for the conduct of the final phase of its investigations on methionine from France, Japan, and Spain following a preliminary determination by the U.S. Department of Commerce (“Commerce”) that imports of methionine from France were being sold at less than fair value (“LTFV”). Notice of the scheduling of the final phase of the Commission’s investigations and of a public hearing to be held in connection therewith was given by posting copies of the notice in the Office of the Secretary, U.S. International Trade Commission, Washington, DC, and by publishing the notice in the Federal Register of March 9, 2021 (86 FR 13585). In light of the restrictions on access to the Commission building due to the COVID–19 pandemic, the Commission conducted its hearing through video conference on May 11, 2021. All persons who requested the opportunity were permitted to participate.

The Commission subsequently issued its final determination that an industry in the United States was materially injured by reason of imports of methionine provided for in subheadings 2930.40.00 and 2930.90.46 of the Harmonized Tariff Schedule of the United States (“HTSUS”) from France that have been found by Commerce to be sold in the United States at LTFV.

Commerce has issued final affirmative antidumping duty determinations with respect to methionine from Japan and Spain. Accordingly, the Commission currently is issuing a supplemental schedule for its antidumping duty investigations on imports of methionine from Japan and Spain.

This supplemental schedule is as follows: The deadline for filing supplemental party comments on Commerce’s final antidumping duty determinations is August 6, 2021. Supplemental party comments may address only Commerce’s final antidumping duty determinations regarding imports of methionine from Japan and Spain. These supplemental final comments may not contain new factual information and may not exceed five (5) pages in length. The supplemental staff report in the final phase of these investigations regarding subject imports from Japan and Spain will be placed in the nonpublic record on August 17, 2021; a public version will be issued thereafter.

For further information concerning these investigations see the Commission’s notice cited above and the Commission’s Rules of Practice and Procedure, part 201, subparts A and B (19 CFR part 201), and part 207, subparts A and C (19 CFR part 207).

Please note the Secretary’s Office will accept only electronic filings during this time. Filings must be made through the Commission’s Electronic Document Information System (EDIS, https://edis.usitc.gov). No in-person paper-based filings or paper copies of any electronic filings will be accepted until further notice.

Additional written submissions to the Commission, including requests pursuant to section 201.12 of the Commission’s rules, shall not be accepted unless good cause is shown for accepting such submissions, or unless the submission is pursuant to a specific request by a Commissioner or Commission staff.

In accordance with sections 201.16(c) and 207.3 of the Commission’s rules, each document filed by a party to the investigations must be served on all other parties to the investigations (as identified by either the public or BPI 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 section 207.21 of the Commission’s rules.

By order of the Commission.


Lisa Barton.
Secretary to the Commission.

[FR Doc. 2021–16375 Filed 7–30–21; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[Investigation Nos. 701–TA–467 and 731–TA–1164–1165 (Second Review)]

Narrow Woven Ribbons With Woven Selvedge From China and Taiwan;
Institution of Five-Year Reviews


ACTION: Notice.

SUMMARY: The Commission hereby gives notice that it has instituted reviews pursuant to the Tariff Act of 1930 (“the Act”), as amended, to determine whether revocation of the countervailing duty order on imports of narrow woven ribbons with woven selvedge (“narrow woven ribbons”) from China and revocation of the antidumping duty orders on narrow woven ribbons from China and Taiwan would be likely to lead to continuation or recurrence of material injury. Pursuant to the Act, interested parties are requested to respond to this notice by submitting the information specified below to the Commission.

DATES: Instituted August 2, 2021. To be assured of consideration, the deadline for responses is September 1, 2021. Comments on the adequacy of responses may be filed with the Commission by October 15, 2021.

FOR FURTHER INFORMATION CONTACT:

General information concerning the Commission may also be obtained by accessing its internet server (https://www.usitc.gov). The public record for this proceeding may be viewed on the Commission’s electronic docket (EDIS) at https://edis.usitc.gov.

SUPPLEMENTARY INFORMATION:

Background.—On September 1, 2010, the Department of Commerce (“Commerce”) issued a countervailing duty order on imports of narrow woven ribbons from China (75 FR 53642) and antidumping duty orders on imports of narrow woven ribbons from China and Taiwan (75 FR 53632, as amended on September 17, 2010, 75 FR 56982).

Following the first five-year reviews by Commerce and the Commission,
effective September 22, 2016, Commerce
issued a continuation of the
countervailing duty order for China and
antidumping duty orders for China and
Taiwan on imports of narrow woven
ribbons (81 FR 65341). The Commission
is now conducting a second review
pursuant to section 751(c) of the Act, as
amended (19 U.S.C. 1675(c)), to
determine whether revocation of the
orders would be likely to lead to
continuation or recurrence of material
injury to the domestic industry within
a reasonably foreseeable time.
Provisions concerning the conduct of
this proceeding may be found in the
Commission’s Rules of Practice and
Procedure at 19 CFR part 201, subparts
A and B, and 19 CFR part 207, subparts
A and F. The Commission will assess
the adequacy of interested party
responses to this notice of institution
to determine whether to conduct full
reviews or expedited reviews. The
Commission’s determination in any
expedited review will be based on the
facts available, which may include
information provided in response to this
notice.
Definitions.—The following
definitions apply to this review:
(1) Subject Merchandise is the class or
kind of merchandise that is within the
scope of the five-year review, as defined by the Department of Commerce.
(2) The Subject Countries in these
reviews are China and Taiwan.
(3) The Domestic Like Product is the
domestically produced product or
products which are like, or in the
absence of like, most similar in
characteristics and uses with the,
Subject Merchandise. In its original
determinations and first five-year
review determinations, the Commission
defined a single Domestic Like Product
consisting of narrow woven ribbons that
are coextensive with Commerce’s scope.
(4) The Domestic Industry is the U.S.
producers as a whole of the Domestic
Like Product, or those producers whose
collective output of the Domestic Like
Product constitutes a major proportion
of the total domestic production of the
product. In its original determinations
and first five-year review
determinations, the Commission
defined the Domestic Industry as all
U.S. producers of narrow woven
ribbons.
(5) An Importer is any person or firm
engaged, either directly or through a
parent company or subsidiary, in
importing the Subject Merchandise into
the United States from a foreign
manufacturer or through its selling
agent.

Participation in the proceeding and
public service list.—Persons, including
industrial users of the Subject
Merchandise and, if the merchandise is
sold at the retail level, representative
consumer organizations, wishing to
participate in the proceeding as parties
must file an entry of appearance with
the Secretary to the Commission, as
provided in §201.11(b)(4) of the
Commission’s rules, no later than 21
days after publication of this notice in
the Federal Register. The Secretary will
maintain a public service list containing
the names and addresses of all persons,
or their representatives, who are parties
to the proceeding.
Former Commission employees who are
seeking to appear in Commission
five-year reviews are advised that they
may appear in a review even if they
participated personally and
substantially in the corresponding
underlying original investigation or an
earlier review of the same underlying
investigation. The Commission’s
designated agency ethics official has
advised that a five-year review is not the
same particular matter as the underlying
original investigation and a five-year
review is not the same particular matter
as an earlier review of the same
underlying investigation for purposes of
18 U.S.C. 207, the post-employment
statute for Federal employees, and
Commission rule 201.15(b) (19 CFR
201.15(b)), 79 FR 3246 (Jan. 17, 2014),
73 FR 24609 (May 5, 2008).
Consequently, former employees are not
required to seek Commission approval
to appear in a review under Commission
rule 19 CFR 201.15, even if the
commission is underlying original
investigation or an earlier review of the
same underlying investigation was
pending when they were Commission
employees. For further ethics advice on
this matter, contact Charles Smith,
Office of the General Counsel, at 202–
205–3408.

Limited disclosure of business
proprietary information (BPI) under an
administrative protective order (APO)
and APO service list.—Pursuant to
§207.7(a) of the Commission’s rules, the
Secretary will make BPI submitted in
this proceeding available to authorized
applicants under the APO issued in the
proceeding, provided that the
application is made no later than 21
days after publication of this notice in
the Federal Register. Authorized
applicants must represent interested
parties, as defined in 19 U.S.C. 1677(9),
who are parties to the proceeding. A
separate service list will be maintained
by the Secretary for those parties
authorized to receive BPI under the
APO.

Certification.—Pursuant to §207.3 of
the Commission’s rules, any person
submitting information to the
Commission in connection with this
proceeding must certify that the
information is accurate and complete to
the best of the submitter’s knowledge. In
making the certification, the submitter
will acknowledge that information
submitted in response to this request for
information and throughout this
proceeding or other proceeding may be
disclosed to and used: (i) By the
Commission, its employees and Offices,
and contract personnel (a) for
developing or maintaining the records
of this or a related proceeding, or (b) in
internal investigations, audits, reviews,
and evaluations relating to the
programs, personnel, and operations of
the Commission including under 5
U.S.C. Appendix 3; or (ii) by U.S.
government employees and contract
personnel, solely for cybersecurity
purposes. All contract personnel will
sign appropriate nondisclosure
agreements.

Written submissions.—Pursuant to
§207.61 of the Commission’s rules, each
terested party is required to respond to
this notice. Each response must provide
the information specified below. The deadline for filing such
responses is September 1, 2021.
Pursuant to §207.62(b) of the
Commission’s rules, eligible parties (as
specified in Commission rule
207.62(b)(1)) may also file comments
concerning the adequacy of responses to
the notice of institution and whether the
Commission should conduct expedited
or full reviews. The deadline for filing
such comments is October 15, 2021. All
written submissions must conform with
the provisions of §201.8 of the
Commission’s rules; any submissions
that contain BPI must also conform with
the requirements of §§201.6, 207.3, and
207.7 of the Commission’s rules. The
Commission’s Handbook on Filing
Procedures, available on the
Commission’s website at https://
www.usitc.gov/documents/handbook_
on_filing_procedures.pdf, elaborates
upon the Commission’s procedures with
respect to filings. Also, in accordance with §§201.16(c) and 207.3 of the
Commission’s rules, each document
filed by a party to the proceeding must be
served on all other parties to the
proceeding (as identified by either the
public or APO service list as
appropriate), and a certificate of service
must accompany the document (if you are
not a party to the proceeding you do
not need to serve your response).
Please note the Secretary’s Office will
accept only electronic filings at this
time. Filings must be made through the
Commission’s Electronic Document
Information System (EDIS, https://
edis.usitc.gov). No in-person paper-
based filings or paper copies of any electronic filings will be accepted until further notice.

No response to this request for information is required if a currently valid Office of Management and Budget ("OMB") number is not displayed; the OMB number is 3117 0016/USITC No. 21–5–496, expiration date June 30, 2023. Public reporting burden for the request is estimated to average 15 hours per response. Please send comments regarding the accuracy of this burden estimate to the Office of Investigations, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436.

Inability to provide requested information.—Pursuant to §207.61(c) of the Commission's rules, any interested party that cannot furnish the information requested by this notice in the requested form and manner shall notify the Commission at the earliest possible time, provide a full explanation of why it cannot provide the requested information, and indicate alternative forms in which it can provide equivalent information. If an interested party does not provide this notification (or the Commission finds the explanation provided in the notification inadequate) and fails to provide a complete response to this notice, the Commission may take an adverse inference against the party pursuant to §776(b) of the Act (19 U.S.C. 1677e(b)) in making its determinations in the reviews.

Information To Be Provided in Response to This Notice of Institution: If you are a domestic producer, union/worker group, or trade/business association; import/export Subject Merchandise from more than one Subject Country; or produce Subject Merchandise in more than one Subject Country, you may file a single response. If you do so, please ensure that your response to each question includes the information requested for each pertinent Subject Country. As used below, the term “firm” includes any related firms.

(1) The name and address of your firm or entity (including World Wide Web address) and name, telephone number, fax number, and Email address of the certifying official.

(2) A statement indicating whether your firm/entity is an interested party under 19 U.S.C. 1677(9) and if so, how, including whether your firm/entity is a U.S. producer of the Domestic Like Product, a U.S. union or worker group, a U.S. importer of the Subject Merchandise, a foreign producer or exporter of the Subject Merchandise, a U.S. or foreign trade or business association (a majority of whose members are interested parties under the statute), or another interested party (including an explanation). If you are a union/worker group or trade/business association, identify the firms in which your workers are employed or which are members of your association.

(3) A statement indicating whether your firm/entity is willing to participate in this proceeding by providing information requested by the Commission.

(4) A statement of the likely effects of the revocation of the countervailing duty order on imports of narrow woven ribbons from China and revocation of the antidumping duty orders on narrow woven ribbons from China and Taiwan on the Domestic Industry in general and/or your firm/entity specifically. In your response, please discuss the various factors specified in section 752(a) of the Act (19 U.S.C. 1675a(a)) including the likely volume of subject imports, likely price effects of subject imports, and likely impact of imports of Subject Merchandise on the Domestic Industry.

(5) A list of all known and currently operating U.S. producers of the Domestic Like Product. Identify any known related parts and the nature of the relationship as defined in section 771(4)(B) of the Act (19 U.S.C. 1677(4)(B)).

(6) A list of all known and currently operating U.S. importers of the Subject Merchandise and producers of the Subject Merchandise in each Subject Country that currently export or have exported Subject Merchandise to the United States or other countries after 2015.

(7) A list of 3–5 leading purchasers in the U.S. market for the Domestic Like Product and the Subject Merchandise (including street address, World Wide Web address, and the name, telephone number, fax number, and Email address of a responsible official at each firm).

(8) A list of known sources of information on national or regional prices for the Domestic Like Product or the Subject Merchandise in the U.S. or other markets.

(9) If you are a U.S. producer of the Domestic Like Product, provide the following information on your firm’s operations on that product during calendar year 2020 (report quantity data in square yards and value data in U.S. dollars). If you are a trade/business association, provide the information, on an aggregate basis, for the firms which are members of your association.

(a) Production (quantity) and, if known, an estimate of the percentage of total U.S. production of the Domestic Like Product accounted for by your firm(s)’ production;

(b) Capacity (quantity) of your firm to produce the Domestic Like Product (that is, the level of production that your establishment(s) could reasonably have expected to attain during the year, assuming normal operating conditions (using equipment and machinery in place and ready to operate), normal operating levels (hours per week/weeks per year), time for downtime, maintenance, repair, and cleanup, and a typical or representative product mix);

(c) the quantity and value of U.S. commercial shipments of the Domestic Like Product produced in your U.S. plant(s);

(d) the quantity and value of U.S. internal consumption/company transfers of the Domestic Like Product produced in your U.S. plant(s); and

(e) the value of (i) net sales, (ii) cost of goods sold (COGS), (iii) gross profit, (iv) selling, general and administrative (SG&A) expenses, and (v) operating income of the Domestic Like Product produced in your U.S. plant(s) (include both U.S. and export commercial sales, internal consumption, and company transfers) for your most recently completed fiscal year (identify the date on which your fiscal year ends).

(10) If you are a U.S. importer or a trade/business association of U.S. importers of the Subject Merchandise from any Subject Country, provide the following information on your firm(s)’ operations on that product during calendar year 2020 (report quantity data in square yards and value data in U.S. dollars). If you are a trade/business association, provide the information, on an aggregate basis, for the firms which are members of your association.

(a) The quantity and value (landed, duty-paid but not including antidumping or countervailing duties) of U.S. imports and, if known, an estimate of the percentage of total U.S. imports of Subject Merchandise from each Subject Country accounted for by your firm(s)’ imports;

(b) the quantity and value (f.o.b. U.S. port, including antidumping and/or countervailing duties) of U.S. commercial shipments of Subject Merchandise imported from each Subject Country;

(c) the quantity and value (f.o.b. U.S. port, including antidumping and/or countervailing duties) of U.S. internal consumption/company transfers of Subject Merchandise imported from each Subject Country.
Merchandise producers or exporters of the Subject Merchandise in any Subject Country, provide the following information on your firm’s(s’) operations on that product during calendar year 2020 (report quantity data in square yards and value data in U.S. dollars, landed and duty-paid at the U.S. port but not including antidumping or countervailing duties). If you are a trade/business association, provide the information, on an aggregate basis, for the firms which are members of your association.

(a) Production (quantity) and, if known, an estimate of the percentage of total production of Subject Merchandise in each Subject Country accounted for by your firm’s(s’) production;

(b) Capacity (quantity) of your firm(s) to produce the Subject Merchandise in each Subject Country (that is, the level of production that your establishment(s) could reasonably have expected to attain during the year, assuming normal operating conditions (using equipment and machinery in place and ready to operate), normal operating levels (hours per week/weeks per year), time for downtime, maintenance, repair, and cleanup, and a typical or representative product mix); and

c) the quantity and value of your firm’s(s’) exports to the United States of Subject Merchandise and, if known, an estimate of the percentage of total exports to the United States of Subject Merchandise from each Subject Country accounted for by your firm’s(s’) exports.

(12) Identify significant changes, if any, in the supply and demand conditions or business cycle for the Domestic Like Product that have occurred in the United States or in the market for the Subject Merchandise in each Subject Country after 2015, and significant changes, if any, that are likely to occur within a reasonably foreseeable time. Supply conditions to consider include technology, production methods; development efforts; ability to increase production (including the shift of production facilities used for other products and the use, cost, or availability of major inputs into production); and factors related to the ability to shift supply among different national markets (including barriers to importation in foreign markets or changes in market demand abroad). Demand conditions to consider include end uses and applications; the existence and availability of substitute products; and the level of competition among the Domestic Like Product produced in the United States. Subject Merchandise produced in each Subject Country, and such merchandise from other countries.

(13) OPTIONAL A statement of whether you agree with the above definitions of the Domestic Like Product and Domestic Industry; if you disagree with either or both of these definitions, please explain why and provide alternative definitions.

Authority: This proceeding is being conducted under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to § 207.61 of the Commission’s rules.

By order of the Commission.

Issued: July 26, 2021.

Lisa Barton,
Secretary to the Commission.

The company plans to manufacture the above controlled substances as bulk active pharmaceutical ingredients (API) for use in product development and for distribution to its customers. In reference to drug codes 7360 (Marihuana), and 7370 (Tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetic. No other activities for these drug codes are authorized for this registration.

William T. McDermott,
Assistant Administrator.

DEPARTMENT OF JUSTICE

[DOCKET NO. DEA–877]

BULK MANUFACTURER OF CONTROLLED SUBSTANCES APPLICATION: AMRI Rensselaer, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: AMRI Rensselaer, Inc., has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to Supplemental Information listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before October 1, 2021. Such persons may also file a written request for a hearing on the application on or before October 1, 2021.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on May 31, 2021, AMRI Rensselaer Inc., 33 Riverside Avenue, Rensselaer, New York 12144–2951, 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>Marihuana</td>
<td>7360</td>
<td>I</td>
</tr>
<tr>
<td>Tetrahydrocannabinols</td>
<td>7370</td>
<td>I</td>
</tr>
<tr>
<td>Amphetamine</td>
<td>1100</td>
<td>II</td>
</tr>
<tr>
<td>Lisdexamfetamine</td>
<td>1205</td>
<td>II</td>
</tr>
<tr>
<td>Pentobarbital</td>
<td>2270</td>
<td>II</td>
</tr>
<tr>
<td>ANPP (4-Anilino-N-phenethyl-4-piperidinamide)</td>
<td>8333</td>
<td>II</td>
</tr>
<tr>
<td>Codeine</td>
<td>9050</td>
<td>II</td>
</tr>
<tr>
<td>Oxycodone</td>
<td>9143</td>
<td>II</td>
</tr>
<tr>
<td>Hydromorphone</td>
<td>9150</td>
<td>II</td>
</tr>
<tr>
<td>Hydrocodone</td>
<td>9193</td>
<td>II</td>
</tr>
<tr>
<td>Morphine</td>
<td>9300</td>
<td>II</td>
</tr>
<tr>
<td>Fentanyl</td>
<td>9801</td>
<td>II</td>
</tr>
</tbody>
</table>

The company plans to manufacture the above controlled substances as bulk active pharmaceutical ingredients (API) for use in product development and for distribution to its customers. In reference to drug codes 7360 (Marihuana), and 7370 (Tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetic. No other activities for these drug codes are authorized for this registration.

William T. McDermott,
Assistant Administrator.

DEPARTMENT OF JUSTICE

[OMB NUMBER 1122–0027]

AGENCY INFORMATION COLLECTION ACTIVITIES; PROPOSED ECOLLECTION REQUESTED; EXTENSION OF A CURRENTLY APPROVED COLLECTION

AGENCY: Office on Violence Against Women, Department of Justice.

ACTION: 60-Day notice.

SUMMARY: The Office on Violence Against Women (OVW), Department of Justice, will be submitting the following information collection request to the Office of Management and Budget (OMB) 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 October 1, 2021.

FOR FURTHER INFORMATION CONTACT: Written comments and/or suggestion regarding the items contained in this notice, especially the estimated public burden and associated response time, should be directed to Cathy Poston, Office on Violence Against Women, at 202–514–5430 or Catherine.poston@usdoj.gov.

SUPPLEMENTARY INFORMATION: Written comments and suggestions from the