TEXAS
Bexar County
Aurora Apartment Hotel, 509 Howard St., San Antonio, SG100006722

Additional documentation has been received for the following resources:

KENTUCKY
Jefferson County
St. James-Belgravia Historic District
(Additional Documentation), Roughly bounded by Central Park, South 4th, South 6th, and Hill Sts., Louisville, AD72005538
Old Louisville Residential District
(Additional Documentation), Irregular pattern roughly bounded by South 7th St., North-South Expwy., Kentucky St., and Avery St., Louisville, AD75000772

Nomination submitted by Federal Preservation Officer:
The State Historic Preservation Officer reviewed the following nomination and responded to the Federal Preservation Officer within 45 days of receipt of the nomination and supports listing the property in the National Register of Historic Places.

ARKANSAS
Newton County
Henderson, Frank and Eva Barnes “Granny,”
Farm, Southwest of Hemmed In Hollow, approx. 1/10 mi. west of Buffalo R. just south of Sneds Cz., Compton vicinity, SG100006726

Authority: Section 60.13 of 36 CFR part 60
Dated: June 8, 2021.

Sherry A. Frear,
Chief, National Register of Historic Places/National Historic Landmarks Program.

[FR Doc. 2021–12837 Filed 6–16–21; 8:45 am]
BILLING CODE 4312–52–P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 731–TA–1528 (Final)]

Seamless Refined Copper Pipe and Tube From Vietnam; Cancellation of Hearing for a Final Phase Anti-Dumping Duty Investigation


ACTION: Notice.

DATES: June 11, 2021.

FOR FURTHER INFORMATION CONTACT:

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.

INTERNATIONAL TRADE COMMISSION

[Investigation No. AA1921–167 (Fifth Review)]

Pressure Sensitive Plastic Tape From Italy; Termination of Five-Year Review


ACTION: Notice.

SUMMARY: The Commission instituted the subject five-year review in March 1, 2021 to determine whether revocation of the antidumping duty order on pressure sensitive plastic tape from Italy would be likely to lead to continuation or recurrence of material injury. On June 7, 2021, the Department of Commerce published notice that it was revoking the order effective April 14, 2021, because no domestic interested party filed a timely notice of intent to participate. Accordingly, the subject review is terminated.

DATES: April 14, 2021 (effective date of revocation of the order).

FOR FURTHER INFORMATION CONTACT:
Andres Andrade (202–205–2078), 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).

Authority: This review is being terminated under authority of title VII of the Tariff Act of 1930 and pursuant to section 751(c) of the Tariff Act of 1930 (19 U.S.C. 1675(c)). This notice is published pursuant to § 207.69 of the Commission’s rules (19 CFR 207.69).

By order of the Commission.
Issued: June 11, 2021.

Katherine Hiner,
Acting Secretary to the Commission.

[FR Doc. 2021–12730 Filed 6–16–21; 8:45 am]
BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

Notice of Receipt of Complaint; Solicitation of Comments Relating to the Public Interest


For an extended description of this document, please refer to the Federal Register.
ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission has received a complaint entitled Certain Silicon Photovoltaic Cells and Modules with Nanostructures, and Products Containing the Same, DN 3552; the Commission is soliciting comments on any public interest issues raised by the complaint or complainant’s filing pursuant to the Commission’s Rules of Practice and Procedure.


General information concerning the Commission may also be obtained by accessing its internet server at United States International Trade Commission (USITC) at https://www.usitc.gov. The public record for this investigation may be viewed on the Commission’s Electronic Document Information System (EDIS) at https://edis.usitc.gov. Hearing-impaired persons are advised that information on this matter can be obtained by contacting the Commission’s TDD terminal on (202) 205–1810.

SUPPLEMENTARY INFORMATION: The Commission has received a complaint and a submission pursuant to § 210.8(b) of the Commission’s Rules of Practice and Procedure filed on behalf of Advanced Silicon Group Technologies, LLC on June 11, 2021. The complaint alleges violations of section 337 of the Tariff Act of 1930 (19 U.S.C. 1337) in the importation into the United States, the sale for importation, and the sale within the United States after importation of certain silicon photovoltaic cells and modules and nanostructures, and products containing the same. The complainant names as respondents: Canadian Solar, Inc. of Canada; Canadian Solar International Limited of China; Canadian Solar Manufacturing (Changshu) Co. Inc. of China; Canadian Solar Manufacturing (Luoyang) Inc. of China; Canadian Solar Manufacturing (Thailand) Co. Ltd. of Thailand; Canadian Solar Manufacturing Vietnam Co., Ltd. of Vietnam; Canadian Solar Solutions, Inc. of Canada; Canadian Solar Construction (USA) LLC of Walnut Creek, CA; Canadian Solar (USA) Inc. of Walnut Creek, CA; Recurrent Energy Group, Inc. of San Francisco, CA; Recurrent Energy LLC of Walnut Creek, CA; Recurrent Energy SH Proco LLC of Walnut Creek, CA; Hanwha Q CELLS & Advanced Materials Corp of South Korea; Hanwha Q Cells GmbH of Germany; Hanwha Q Cells Malaysia Sdn. Bhd. of Malaysia; Hanwha Q Cells (Qidong) Co., Ltd. of China; Hanwha Solutions Corporation of South Korea; Hanwha Energy USA Holding Corp. (d/b/a 174 Power Global Corporation) of Irvine, CA; Hanwha Q Cell EPC USA LLC of Irvine, CA; Hanwha Q Cells America Inc. of Irvine, CA; Hanwha Q Cells USA Corp. of Irvine, CA; Hanwha Q Cells USA Inc. of Dalton, GA; HQC Rock River Solar Holdings LLC of Irvine, CA; HQC Rock River Solar Power Generation Station, LLC of Beloit, WI; Boviet Solar Technology Co., Ltd. of Vietnam; Ningbo Boway Alloy Material Co., Ltd. of China; Boviet Renewable Power LLC of San Jose, CA; and Boviet Solar USA Ltd. of San Jose, CA. The complainant requests that the Commission issue a limited exclusion order; cease and desist orders, and impose a bond upon respondent alleged infringing articles during the 60-day Presidential review period pursuant to 19 U.S.C. 1337(j).

Proposed respondents, other interested parties, and members of the public are invited to file comments on any public interest issues raised by the complaint or § 210.8(b) filing. Comments should address whether issuance of the relief specifically requested by the complainant in this investigation would affect the public health and welfare in the United States, competitive conditions in the United States economy, the production of like or directly competitive articles in the United States, or United States consumers.

In particular, the Commission is interested in comments that:
(i) Explain how the articles potentially subject to the requested remedial orders are used in the United States;
(ii) identify any public health, safety, or welfare concerns in the United States related to the requested remedial orders;
(iii) identify like or directly competitive articles that complainant, its licensees, or third parties make in the United States which could replace the subject articles if they were to be excluded;
(iv) indicate whether complainant, complainant’s licensees, and/or third party suppliers have the capacity to replace the volume of articles potentially subject to the requested exclusion order and/or a cease and
desist order within a commercially reasonable time; and
(v) explain how the requested remedial orders would impact United States consumers.

Written submissions on the public interest must be filed no later than by close of business, eight calendar days after the date of publication of this notice in the Federal Register. There will be further opportunities for comment on the public interest after the issuance of any final initial determination in this investigation. Any written submissions on other issues must also be filed by no later than the close of business, eight calendar days after publication of this notice in the Federal Register. Complainant may file replies to any written submissions no later than three calendar days after the date on which any initial submissions were due. No other submissions will be accepted, unless requested by the Commission. Any submissions and replies filed in response to this Notice are limited to five (5) pages in length, inclusive of attachments.

Persons filing written submissions must file the original document electronically on or before the deadlines stated above. Submissions should refer to the docket number (“Docket No. 3552”) in a prominent place on the cover page and/or the first page. (See Handbook for Electronic Filing Procedures, Electronic Filing Procedures). 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. Persons with questions regarding filing should contact the Secretary at EDIS3Help@usitc.gov.

Any person desiring to submit a document to the Commission in confidence must request confidential treatment. All such requests should be directed to the Secretary to the Commission and must include a full statement of the reasons why the Commission should grant such treatment. See 19 CFR 201.6. Documents for which confidential treatment by the Commission is properly sought will be treated accordingly. All information, including confidential business information and documents for which confidential treatment is properly sought, submitted to the Commission for

1 Handbook for Electronic Filing Procedures:
purposes of this Investigation 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 nonconfidential written submissions will be available for public inspection at the Office of the Secretary and on EDIS.3

This action is taken under the authority of section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and of §§ 201.10 and 210.8(c) of the Commission’s Rules of Practice and Procedure (19 CFR 201.10, 210.8(c)). By order of the Commission. Issued: June 11, 2021.

Lisa Barton, Secretary to the Commission.

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–854]

Importer of Controlled Substances Application: Xcelience

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

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

<table>
<thead>
<tr>
<th>Controlled substance</th>
<th>Drug code</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>2-(4-Iodo-2,5-dimethoxyphenyl)ethanamine</td>
<td>7518</td>
<td>I</td>
</tr>
<tr>
<td>2-(2,5-dimethoxyphenyl)ethanamine</td>
<td>7517</td>
<td>I</td>
</tr>
<tr>
<td>3,4-methylenedioxymethamphetamine</td>
<td>7405</td>
<td>I</td>
</tr>
<tr>
<td>3,4-methylenedioxyamphetamine</td>
<td>7400</td>
<td>I</td>
</tr>
<tr>
<td>3,4,5-trimethoxyamphetamine</td>
<td>7390</td>
<td>I</td>
</tr>
</tbody>
</table>

The company plans to import drug code 7437 (Psilocybin) as finished dosage form for clinical trials, research, and analytical purposes. 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.

William T. McDermott, Assistant Administrator.

[FR Doc. 2021–12816 Filed 6–16–21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–855]

Bulk Manufacturer of Controlled Substances Application: Organix Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Organix Inc. has applied to be registered as a bulk manufacturer of the following 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 July 19, 2021. Such persons may also file a written request for a hearing on the application on or before July 19, 2021.

ADDITIONAL INFORMATION: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on April 27, 2021, Organix Inc., 240 Salem Street, Woburn, Massachusetts 01801–2029, 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>Psilocybin</td>
<td>7437</td>
<td>I</td>
</tr>
</tbody>
</table>

The company plans to import drug code 7437 (Psilocybin) as finished dosage form for clinical trials, research, and analytical purposes. 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.

William T. McDermott, Assistant Administrator.

[FR Doc. 2021–12816 Filed 6–16–21; 8:45 am]

BILLING CODE P

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on April 28, 2021, Xcelience, 4901 West Grace Street, Tampa, Florida 33607–3805, applied to be registered as an importer 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>Psilocybin</td>
<td>7437</td>
<td>I</td>
</tr>
</tbody>
</table>

The company plans to import drug code 7437 (Psilocybin) as finished dosage form for clinical trials, research, and analytical purposes. 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.

William T. McDermott, Assistant Administrator.

[FR Doc. 2021–12816 Filed 6–16–21; 8:45 am]

BILLING CODE P