

1, 5, 6, and 12 of the '896 patent. On July 25, 2022, Sonos filed an opposition to Google's modification petition.

The Commission has determined that Google's petition complies with the requirements for institution of a modification proceeding pursuant to Commission Rule 210.76. Accordingly, the Commission has determined to institute a modification proceeding and has delegated the proceeding to the CALJ. The presiding CALJ shall submit a recommended determination within six (6) months after publication of notice of this Order in the **Federal Register**. Sonos, Google, and OUII are named as parties to the modification proceeding.

The Commission vote for this determination took place on August 8, 2022.

The authority for the Commission's determination is contained in section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and in Part 210 of the Commission's Rules of Practice and Procedure (19 CFR part 210).

By order of the Commission.

Issued: August 9, 2022.

Katherine Hiner,

Acting Secretary to the Commission.

[FR Doc. 2022-17398 Filed 8-11-22; 8:45 am]

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INTERNATIONAL TRADE COMMISSION

[Investigation Nos. 731-TA-825-826 (Fourth Review)]

Certain Polyester Staple Fiber From South Korea and Taiwan

Determinations

On the basis of the record¹ developed in the subject five-year reviews, the United States International Trade Commission ("Commission") determines, pursuant to the Tariff Act of 1930 ("the Act"), that revocation of the antidumping duty orders on certain polyester staple fiber from South Korea and Taiwan would be likely to lead to continuation or recurrence of material injury to an industry in the United States within a reasonably foreseeable time.

Background

The Commission instituted these reviews on January 3, 2022 (87 FR 119) and determined on April 8, 2022, that it would conduct expedited reviews (87 FR 38780, June 29, 2022).

¹ The record is defined in § 207.2(f) of the Commission's Rules of Practice and Procedure (19 CFR 207.2(f)).

The Commission made these determinations pursuant to section 751(c) of the Act (19 U.S.C. 1675(c)). It completed and filed its determinations in these reviews on August 8, 2022. The views of the Commission are contained in USITC Publication 5341 (August 2022), entitled *Certain Polyester Staple Fiber from South Korea and Taiwan: Investigation Nos. 731-TA-825-826 (Fourth Review)*.

By order of the Commission.

Issued: August 8, 2022.

Katherine Hiner,

Acting Secretary to the Commission.

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INTERNATIONAL TRADE COMMISSION

[Investigation Nos. 701-TA-540-543 and 731-TA-1283-1287 and 1290 (Review)]

Cold-Rolled Steel Flat Products From Brazil, China, India, Japan, South Korea, and the United Kingdom

Determinations

On the basis of the record¹ developed in the subject five-year reviews, the United States International Trade Commission ("Commission") determines, pursuant to the Tariff Act of 1930 ("the Act"), that revocation of the countervailing duty orders on cold-rolled steel flat products ("cold-rolled steel") from China, India, and South Korea and the antidumping duty orders on cold-rolled steel from China, India, Japan, South Korea, and the United Kingdom would be likely to lead to continuation or recurrence of material injury to an industry in the United States within a reasonably foreseeable time. The Commission further determines that revocation of the countervailing and antidumping duty orders on cold-rolled steel from Brazil would not be likely to lead to continuation or recurrence of material injury to an industry in the United States within a reasonably foreseeable time.²

¹ The record is defined in § 207.2(f) of the Commission's Rules of Practice and Procedure (19 CFR 207.2(f)).

² Commissioners Rhonda K. Schmidlein and Randolph J. Stayin determine that revocation of the countervailing duty orders on CRS from Brazil, China, India, and South Korea and the antidumping duty orders on CRS from Brazil, China, India, Japan, South Korea, and the United Kingdom would be likely to lead to continuation or recurrence of material injury to an industry in the United States within a reasonably foreseeable time.

Background

The Commission instituted these reviews on June 1, 2021 (86 FR 29286) and determined on September 7, 2021 that it would conduct full reviews (86 FR 52180, September 20, 2021). Notice of the scheduling of the Commission's reviews 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** on December 13, 2021 (86 FR 70864). The Commission conducted its hearing on May 24, 2022. All persons who requested the opportunity were permitted to participate.

The Commission made these determinations pursuant to section 751(c) of the Act (19 U.S.C. 1675(c)). It completed and filed its determinations in these reviews on August 9, 2022. The views of the Commission are contained in USITC Publication 5339 (August 2022), entitled *Cold-Rolled Steel Flat Products from Brazil, China, India, Japan, South Korea, and the United Kingdom: Investigation Nos. 701-TA-540-543 and 731-TA-1283-1287 and 1290 (Review)*.

By order of the Commission.

Issued: August 9, 2022.

Katherine Hiner,

Acting Secretary to the Commission.

[FR Doc. 2022-17399 Filed 8-11-22; 8:45 am]

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INTERNATIONAL TRADE COMMISSION

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

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission has received a complaint entitled *Certain Pillows and Seat Cushions, Components Thereof, and Packaging Thereof, DN 3633*; 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.

FOR FURTHER INFORMATION CONTACT:

Katherine M. Hiner, Acting Secretary to the Commission, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436, telephone (202) 205-2000. The public version of the

complaint can be accessed on the Commission's Electronic Document Information System (EDIS) at <https://edis.usitc.gov>. For help accessing EDIS, please email EDIS3Help@usitc.gov. 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 Purple Innovation, LLC on August 5, 2022. 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 pillows and seat cushions, components thereof, and packaging thereof. The complainant names as respondents: Bedmate-U Co., Ltd. of Korea; Chuang Fan Handicraft Co., Ltd. of China; Dongguan Bounce Technology Co., Ltd. of China; Dongguan Jingrui Silicon Technology Co., Ltd. of China; Foshan Dirani Design Furniture Co., Ltd. of China; Global Ocean Trading Co., Ltd. of China; Guang An Shi Lin Chen Zai Sheng Wuzi Co. Ltd. of China; Guang Zhou Wen Jie Shang Mao Youxian Gongsi Co., Ltd. of China; Guangzhou Epsilon Import and Export Co., Ltd. of China; Guangzhoushi Baixiangguo Keji Youxian Gongsi Co., Ltd. of China; Haircrafters LLC of Chattanooga, TN; Hangzhou Lishang Import & Export Co., Ltd. of China; Hangzhou Lydia Sports Goods Co., Ltd. of China; Hebei Zeyong Technology Co., Ltd. of China; Henson Holdings, LLC d.b.a. SelectSoma of Lafayette, LA; Hetaibao of China; Hubei Sheng Bingyi Dianzi Keji Youxian Gongsi Co. Ltd. of China; Kaifeng Shi Long Ting Qu Chen Yi Shangmao Youxian Gongsi Co., Ltd. of China; Lankao Junchang Electronic Commerce Co., Ltd. of China; Lei Lei Wang of China; Liu Lin Xian Xu Bin Dian Zi Chan Pin Dian of China; Nanchang Shirong Bao Er Guanggao Youxian Gongsi Co., Ltd. of China; Ningbo Bolian Import & Export Co., Ltd. of China; Ningbo Minzhou Import & Export Co., Ltd. of China; Ruian Ketai Commodity Co. Ltd. of China; Ruian Xiu

Yuan Guoji MaoYi Youxian Gongsi Co., Ltd. of China; Shandong Jiu Hui Xinxi Keji Youxian Gongsi Co., Ltd. of China; Shanxi Chao Ma Xun Keji Youxian Gongsi Co., Ltd. of China; Shenzhen Baibaikang Technology Co., Ltd. of China; Shenzhen Leadfar Industry Co., Ltd. of China; Shenzhen Shi Chi Yang Wang Luo Ji Shu Youxian Gongsi Co., Ltd. of China; Shenzhen Shi Mai Rui Ke Dianzi Shangwu Co. Ltd. of China; Shenzhen Shi Xin Shangpin Dianzi Shangwu Youxian Gongsi Co., Ltd. of China; Shenzhen Shi Yan Huang Chu Hai Keji Youxian Gongsi Co., Ltd. of China; Shenzhen Shi Yuxiang Meirong Yongju Youxian Gongsi Co. Ltd. of China; Shenzhen Tianrun Material Co., Ltd. of China; Wuhan Chenkuxuan Technology Co., Ltd. of China; Xiao Dawei of China; Xiao Xiao Pi Fa Shang Mao You Xian Ze Ren Gongsi Co. of China; YaRu Wang of China; Yiwu Youru E-commerce Co., Ltd. of China; Zhejiang Xinhui Import & Export Co., Ltd. of China; and Zhou Meng Bo of China. The complainant requests that the Commission issue a limited exclusion order and cease and desist orders and impose a bond upon respondents' 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 relating 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. 3633") in a prominent place on the cover page and/or the first page. (See Handbook for Electronic Filing Procedures, Electronic Filing Procedures¹). 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: https://www.usitc.gov/documents/handbook_on_filing_procedures.pdf.

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,² 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: August 8, 2022.

Katherine Hiner,

Acting Secretary to the Commission.

[FR Doc. 2022-17298 Filed 8-11-22; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1067]

Importer of Controlled Substances

Application: Cambridge Isotope Laboratories, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Cambridge Isotope Laboratories has applied to be registered as an importer 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 September 12, 2022. Such persons may also file a written request for a hearing on the application on or before September 12, 2022.

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. All requests for a hearing must 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. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on July 18, 2022, Cambridge Isotope Laboratories, 50 Frontage Road, Andover, Massachusetts 01810-5413, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid.	2010	I
Tetrahydrocannabinols ..	7370	I
Morphine	9300	II

The company plans to import the listed controlled substances for preparation of analytical standards and formulations. In reference to drug codes 7370 (Tetrahydrocannabinols), the company plans to import a synthetic Tetrahydrocannabinol. No other activities for these drug codes are authorized for this registration.

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.

Kristi O'Malley,

Assistant Administrator.

[FR Doc. 2022-17364 Filed 8-11-22; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1065]

Importer of Controlled Substances

Application: Galephar Pharmaceutical Research, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Galephar Pharmaceutical Research, Inc. has applied to be registered as an importer 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 September 12, 2022. Such persons may also file a written request for a hearing on the application on or before September 12, 2022.

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. All requests for a hearing must 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. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on July 28, 2022, Galephar Pharmaceutical Research, Inc., 100 Carr 198 Industrial Park, Juncos, Puerto Rico 00777-3873, applied to be registered as

² All contract personnel will sign appropriate nondisclosure agreements.

³ Electronic Document Information System (EDIS): <https://edis.usitc.gov>.