INTERNATIONAL TRADE COMMISSION

[Investigation Nos. 731–TA–1632, 1634, 1635, and 1639 (Final)]

Mattresses From India, Kosovo, Mexico, and Spain; Determinations

On the basis of the record [1] developed in the subject investigations, the United States International Trade Commission (''Commission'') determines, pursuant to the Tariff Act of 1930 (''the Act''), that an industry in the United States is materially injured by reason of imports of mattresses from India, Kosovo, Mexico, and Spain, provided for in subheadings 9404.21.00, 9404.29.10, and 9404.29.90 of the Harmonized Tariff Schedule of the United States, that have been found by the U.S. Department of Commerce (''Commerce'') to be sold in the United States at less than fair value (''LTFV'').[2]

Background

The Commission instituted these investigations effective July 28, 2023, following receipt of petitions filed with the Commission and Commerce by Brooklyn Bedding LLC, Phoenix, Arizona; Carpenter Company, Richmond, Virginia; Corsicana Mattress Company, Dallas, Texas; Future Foam, Inc., Council Bluffs, Iowa; FXI, Inc., Radnor, Pennsylvania; Kolcraft Enterprises, Inc., Chicago, Illinois; Leggett & Platt, Incorporated, Carthage, Missouri; Serta Simmons Bedding, Inc., Doraville, Georgia; Southerland Inc., Antioch, Tennessee; Tempur Sealy International, Inc., Lexington, Kentucky; the International Brotherhood of Teamsters, Washington, DC; and the United Steel, Paper and Forestry, Rubber, Manufacturing, Energy, Allied Industrial and Service Workers International Union, AFL–CIO, Washington, DC. The Commission scheduled the final phase of the investigations following notification of preliminary determinations by Commerce that imports of mattresses from Bosnia and Herzegovina, Bulgaria, Burma, India, Italy, Kosovo, Mexico, Philippines, Poland, Slovenia, Spain, and Taiwan were being sold at LTFV within the meaning of § 733(b) of the Act (19 U.S.C. 1673b(b)). 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 6, 2024 (89 FR 16026). The Commission conducted its hearing on May 9, 2024. All persons who requested the opportunity were permitted to participate.

Although the antidumping duty petitions for mattresses from Bosnia and Herzegovina, Bulgaria, Burma, India, Italy, Kosovo, Mexico, Philippines, Poland, Slovenia, Spain, and Taiwan were filed on the same day, July 28, 2023, the investigation schedules became staggered when Commerce did not align its investigations concerning Bosnia and Herzegovina, Bulgaria, Burma, Italy, Philippines, Poland, Slovenia, and Taiwan with its investigations concerning India, Kosovo, Mexico, and Spain. On June 28, 2024, the Commission issued final affirmative determinations in its antidumping duty investigations of mattresses from Bosnia and Herzegovina, Bulgaria, Burma, Italy, Philippines, Poland, Slovenia, and Taiwan (89 FR 55657, July 5, 2024). Following notification of final determinations by Commerce that imports of mattresses from India, Kosovo, Mexico, and Spain were being sold at LTFV within the meaning of section 735(a) of the Act (19 U.S.C. 1673d(a)), notice of the supplemental scheduling of the final phase of the Commission's antidumping duty investigations concerning India, Kosovo, Mexico, and Spain 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** (89 FR 60658, July 26, 2024).[3]

The Commission made these determinations pursuant to § 735(b) of the Act (19 U.S.C. 1673d(b)). It completed and filed its determinations in these investigations on August 28, 2024. The views of the Commission are contained in USITC Publication 5539 (August 2024), entitled *Mattresses from India, Kosovo, Mexico, and Spain: Investigation Nos. 731–TA–1632, 1634, 1635, and 1639 (Final).*

By order of the Commission.

Issued: August 28, 2024.

Sharon Bellamy,

Supervisory Hearings and Information Officer.

[FR Doc. 2024–19781 Filed 9–3–24; 8:45 am]

BILLING CODE 7020–02–P

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

[2] 89 FR 59047 (India), 89 FR 59043 (Kosovo), 89 FR 59062 (Mexico), and 89 FR 59059 (Spain), July 22, 2024.

[3] A countervailing duty petition on mattresses from Indonesia was also filed on the same day as the antidumping duty petitions concerning mattresses from Bosnia and Herzegovina, Bulgaria, Burma, India, Italy, Kosovo, Mexico, Philippines, Poland, Slovenia, Spain, and Taiwan. However, Commerce published a final negative countervailing duty determination with respect to mattresses from Indonesia on July 22, 2024 (89 FR 59050). The Commission therefore terminated its countervailing duty investigation on mattresses from Indonesia (89 FR 60661, July 26, 2024).

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1412]

Importer of Controlled Substances Application: Chattem Chemicals

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Chattem Chemicals 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 October 4, 2024. Such persons may also file a written request for a hearing on the application on or before October 4, 2024.

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 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on July 19, 2024, Chattem Chemicals, 3801 Saint Elmo Avenue, Chattanooga, Tennessee 37409–1237, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Methamphetamine	1105	II
4-Anilino-N-phenethyl-4-piperidine (ANPP)	8333	II
Phenylacetone	8501	II
Cocaine	9041	II
Poppy Straw Concentrate	9670	II
Tapentadol	9780	II

The company plans to import the listed controlled substances to manufacture bulk controlled substances for sale to its customers. The company plans to import an intermediate of Tapentadol (9780), to bulk manufacture Tapentadol for distribution to its customers. 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.

Marsha L. Ikner,

Acting Deputy Assistant Administrator.

[FR Doc. 2024–19789 Filed 9–3–24; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1415]

Bulk Manufacturer of Controlled Substances Application: Bright Green Corporation

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Bright Green Corporation 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 November 4, 2024. Such persons may also file a written request for a hearing on the application on or before November 4, 2024.

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 July 22, 2024, Bright Green Corporation, 1033 George Hanosh Boulevard, Grants, New Mexico 87020, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Marihuana Extract	7350	I
Marihuana	7360	I
Tetrahydrocannabinols	7370	I

The company plans to bulk manufacture the listed controlled substances for research purposes. No other activities for these drug codes are authorized for this registration.

Marsha L. Ikner,

Acting Deputy Assistant Administrator.

[FR Doc. 2024–19777 Filed 9–3–24; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1401]

Bulk Manufacturer of Controlled Substances Application: Continuus Pharmaceuticals

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Continuus Pharmaceuticals 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 November 4, 2024. Such persons may also file a written request for a hearing on the application on or before November 4, 2024.

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