must offer you the opportunity to appeal the decision in accordance with the agency's procedures. If the agency has received and acted upon or will act upon a SAVE verification and you do not believe the response is correct, you may make an appointment for an in-person interview at a local USCIS office. Detailed information on how to make corrections or update your immigration record, make an appointment, or submit a written request to correct records under the Freedom of Information Act can be found on the SAVE website at *www.uscis.gov/save.*

Tracy L. Renaud,

Senior Official Performing the Duties of the Director, U.S. Citizenship and Immigration Services.

[FR Doc. 2021–03149 Filed 2–12–21; 8:45 am]

BILLING CODE P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 337–TA–1226]

Certain Artificial Eyelash Extension Systems, Products, and Components Thereof; Commission Determination Not To Review an Initial Determination Granting Complainant's Motion for Leave To Amend the Complaint and Notice of Investigation

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission has determined not to review an initial determination (''ID'') (Order No. 10) of the presiding administrative law judge (''ALJ'') granting the complainant's motion for leave to amend the complaint and notice of investigation.

FOR FURTHER INFORMATION CONTACT: Lynde Herzbach, Office of the General Counsel, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436, telephone (202) 205–3228. Copies of non-confidential documents filed in connection with this investigation may be viewed on the Commission's electronic docket (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 *https://www.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: On October 28, 2020, the Commission instituted this investigation under section 337 of the Tariff Act of 1930, as amended, 19 U.S.C. 1337 (''section 337''), based on a complaint filed by Lashify, Inc. of Glendale, California (''Complainant''). *See* 85 FR 68366–67. The complaint, as supplemented, alleges a violation of section 337 based upon the importation into the United States, sale for importation, or sale after importation into the United States of certain artificial eyelash extension systems, products, and components thereof by reason of infringement of certain claims of U.S. Patent Nos. 10,660,388 and 10,721,984 and U.S. Design Patent Nos. D877,416 and D867,664. The complaint also alleges the existence of a domestic industry. The notice of investigation names nine respondents, including CVS Health Corporation of Woonsocket, Rhode Island and Ulta Beauty, Inc. of Bolingbrook, Illinois. *See id.* The Office of Unfair Import Investigations is also a party to the investigation. *See id.*

On January 8, 2021, Complainant filed a motion seeking leave to amend the complaint and notice of investigation to substitute: (1) CVS Pharmacy, Inc. in place of CVS Health Corporation and (2) Ulta Salon, Cosmetics & Fragrance, Inc. in place of Ulta Beauty, Inc. No responses to the motion were filed.

On January 22, 2021, the ALJ issued the subject ID (Order No. 10) granting Complainant's motion for leave to amend the complaint and notice of investigation to reflect the substitutions. Order No. 10 (Jan. 22, 2021). The subject ID finds that Complainant's motion is supported by good cause pursuant to Commission Rule 210.14(b) (19 CFR 210.14(b)) and that there is no prejudice if the motion is granted. No party petitioned for review of the subject ID.

The Commission has determined not to review the subject ID. CVS Pharmacy, Inc. and Ulta Salon, Cosmetics & Fragrance, Inc. are named as respondents in this investigation; and CVS Health Corporation and Ulta Beauty, Inc. are terminated from the investigation.

The Commission vote for this determination took place on February 10, 2021.

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: February 10, 2021.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2021–03059 Filed 2–12–21; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 731–TA–1059 (Third Review)]

Hand Trucks and Certain Parts Thereof From China

Determination

On the basis of the record [1] developed in the subject five-year review, the United States International Trade Commission (''Commission'') determines, pursuant to the Tariff Act of 1930 (''the Act''), that revocation of the antidumping duty order on hand trucks and certain parts thereof from China 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 this review on July 1, 2020 (85 FR 39584) and determined on October 5, 2020 that it would conduct an expedited review (86 FR 2001, January 11, 2021).

The Commission made this determination pursuant to section 751(c) of the Act (19 U.S.C. 1675(c)). It completed and filed its determination in this review on February 9, 2021. The views of the Commission are contained in USITC Publication 5159 (February 2021), entitled *Hand Trucks and Certain Parts Thereof from China: Investigation No. 731–TA–1059 (Third Review).*

By order of the Commission.

Issued: February 9, 2021.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2021–02975 Filed 2–12–21; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–784]

Importer of Controlled Substances Application: S and B Pharma, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

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

SUMMARY: S&B Pharma, 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 file written comments on or objections to the issuance of the proposed registration on or before March 18, 2021. Such persons may also file a written request for a hearing on the application on or before March 18, 2021.

ADDRESSES: 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: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also 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.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on December 18, 2020, S&B Pharma, Inc., 405 S Motor Avenue, Azusa, California 91702–3232, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
4-Anilino-N-Phenethyl-4-Piperdine (ANPP).	8333	II
Tapentadol	9780	II

The company plans to import intermediate forms of Tapentadol (9780) for further manufacturing prior to distribution to its customers. The company plans to import ANPP (8333) to bulk manufacture other controlled substances for distribution to its customers. No other activity for this drug code is 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.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2021–02979 Filed 2–12–21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–780]

Bulk Manufacturer of Controlled Substances Application: Bulk Manufacturer of Marihuana: North Star Holdings California, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: The Drug Enforcement Administration (DEA) is providing notice of an application it has received from an entity applying to be registered to manufacture in bulk basic class(es) of controlled substances listed in schedule I. DEA intends to evaluate this and other pending applications according to its regulations governing the program of growing marihuana for scientific and medical research under DEA registration.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefor, may file written comments on or objections to the issuance of the proposed registration on or before April 19, 2021.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. To ensure proper handling of comments, please reference Docket No. DEA–780 in all correspondence, including attachments.

SUPPLEMENTARY INFORMATION: The Controlled Substances Act (CSA) prohibits the cultivation and distribution of marihuana except by persons who are registered under the CSA to do so for lawful purposes. In accordance with the purposes specified in 21 CFR 1301.33(a), DEA is providing notice that the entity identified below has applied for registration as a bulk manufacturer of schedule I controlled substances. In response, registered bulk manufacturers of the affected basic class(es), and applicants therefor, may file written comments on or objections of the requested registration, as provided in this notice. This notice does not constitute any evaluation or determination of the merits of the application submitted.

The applicant plans to manufacture bulk active pharmaceutical ingredients (APIs) for product development and distribution to DEA registered researchers. If the application for registration is granted, the registrant would not be authorized to conduct other activity under this registration aside from those coincident activities specifically authorized by DEA regulations. DEA will evaluate the application for registration as a bulk manufacturer for compliance with all applicable laws, treaties, and regulations and to ensure adequate safeguards against diversion are in place.

As this applicant has applied to become registered as a bulk manufacturer of marihuana, the application will be evaluated under the criteria of 21 U.S.C. 823(a). DEA will conduct this evaluation in the manner described in the rule published at 85 FR 82333 on December 18, 2020, and reflected in DEA regulations at 21 CFR part 1318.

In accordance with 21 CFR 1301.33(a), DEA is providing notice that on January 15, 2021, North Star Holdings California, LLC, 69375 Ramon Road, Cathedral City, California 92234, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substances:

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

William T. McDermott,

Assistant Administrator.

[FR Doc. 2021–02980 Filed 2–12–21; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–778]

Importer of Controlled Substances Application: Noramco Coventry, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Noramco Coventry, LLC 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.