

submissions will be available for public inspection on EDIS.

This action is taken under the authority of 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: December 3, 2021.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2021-26626 Filed 12-8-21; 8:45 am]

BILLING CODE 7020-02-P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 701-TA-665 (Final)]

Certain Mobile Access Equipment and Subassemblies Thereof From China; Determination

On the basis of the record¹ developed in the subject investigation, 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 threatened with material injury by reason of imports of certain mobile access equipment and subassemblies thereof ("mobile access equipment") from China, provided for in subheadings 8427.10.80, 8427.20.80, 8427.90.00, and 8431.20.00 of the Harmonized Tariff Schedule of the United States, that have been found by the U.S. Department of Commerce ("Commerce") to be subsidized by the government of China.²

Background

The Commission instituted this investigation effective February 26, 2021, following receipt of a petition filed with the Commission and Commerce by the Coalition of American Manufacturers of Mobile Access Equipment ("CAMMAE" or "the Coalition").³ The Commission scheduled the final phase of the investigation following notification of a preliminary determination by Commerce that imports of mobile access equipment from China were being subsidized within the meaning of section 703(b) of the Act (19 U.S.C. 1671b(b)). Notice of the scheduling of the final phase of the Commission's investigation 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 August 12, 2021 (86 FR 44402). In light of the restrictions on access to the Commission building due to the COVID-19 pandemic, the Commission conducted its hearing through written testimony and video conference on October 12, 2021. All persons who requested the opportunity were permitted to participate.

The Commission made this determination pursuant to § 705(b) of the Act (19 U.S.C. 1671d(b)). It completed and filed its determination in this investigation on December 3, 2021. The views of the Commission are contained in USITC Publication 5242 (December 2021), entitled *Certain Mobile Access Equipment and Subassemblies Thereof from China: Investigation No. 701-TA-665 (Final)*.

By order of the Commission.

Issued: December 3, 2021.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2021-26623 Filed 12-8-21; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-938]

Importer of Controlled Substances Application: Catalent Pharma Solutions, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Catalent Pharma Solutions, LLC has applied to be registered as an importer 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 January 10, 2022. Such persons may also file a written request for a hearing on the application on or before January 10, 2022.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must

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

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on September 10, 2021, Catalent Pharma Solutions LLC, 3031 Red Lion Road, Philadelphia, Pennsylvania 19114, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Psilocybin	7437	I
Psilocyn	7438	I

The company plans to import the above controlled substances as finished dosage unit products for clinical trials, research, and analytical activities. No other activity for these drug codes 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.

Brian S. Besser,

Acting Assistant Administrator.

[FR Doc. 2021-26678 Filed 12-8-21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA 937]

Importer of Controlled Substances Application: Fresenius Kabi USA, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Fresenius Kabi USA, LLC has applied to be registered as an importer 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

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

² 86 FR 57809 (October 19, 2021).

³ The Coalition is composed of JLG Industries, Inc. ("JLG"), Hagerstown, Maryland and Terex Corporation ("Terex"), Redmond, Washington.

applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before January 10, 2022. Such persons may also file a written request for a hearing on the application on or before January 10, 2022.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also 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.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on September 8, 2021, Fresenius Kabi USA, LLC, 3159 Staley Road, Grand Island, New York 14072–2028, applied to be registered as an importer of the following basic class of controlled substance:

Controlled substance	Drug code	Schedule
Remifentanyl	9739	II

The company plans to import the listed controlled substances for bulk manufacture. 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.

Brian S. Besser,
Acting Assistant Administrator.
[FR Doc. 2021–26677 Filed 12–8–21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–928]

Bulk Manufacturer 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 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 February 7, 2022. Such persons may also file a written request for a hearing on the application on or before February 7, 2022.

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

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on September 29, 2021, Noramco Coventry LLC, 498 Washington Street, Coventry, Rhode Island 02816, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Marihuana	7360	I
Tetrahydrocannabinols ..	7370	I
Dihydromorphine	9145	I
Methylphenidate	1724	II
Oxycodone	9143	II
Hydromorphone	9150	II
Hydrocodone	9193	II
Levorphanol	9220	II
Morphine	9300	II
Oripavine	9330	II
Thebaine	9333	II
Oxymorphone	9652	II
Noroxymorphone	9668	II
Tapentadol	9780	II

The company plans to bulk manufacture the listed controlled substances for use as intermediates and converted to other controlled substances or for sale to its customers. In reference to drug codes 7360 (Marihuana), and 7370 (Tetrahydrocannabinols), the company plans to bulk manufacture

these drugs as synthetics. No other activities for these drug codes are authorized for this registration.

Brian S. Besser,
Acting Assistant Administrator.
[FR Doc. 2021–26676 Filed 12–8–21; 8:45 am]
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NATIONAL SCIENCE FOUNDATION

Notice of Permits Issued Under the Antarctic Conservation Act of 1978

AGENCY: National Science Foundation.
ACTION: Notice of permits issued.

SUMMARY: The National Science Foundation (NSF) is required to publish notice of permits issued under the Antarctic Conservation Act of 1978. This is the required notice.

FOR FURTHER INFORMATION CONTACT: Polly Penhale, ACA Permit Officer, Office of Polar Programs, National Science Foundation, 2415 Eisenhower Avenue, Alexandria, VA 22314; 703–292–8030; email: ACApermits@nsf.gov.

SUPPLEMENTARY INFORMATION: On November 5, 2021, the National Science Foundation published a notice in the **Federal Register** of permit applications received. The permits were issued on December 6, 2021, to:

1. Henry Wulff, Atlas Ocean Voyages—Permit No. 2022–021
2. Deirdre Dirkman, Vantage Deluxe World Travel—Permit No. 2022–022
3. Tom Russell, Swan Hellenic Antarctic—Permit No. 2022–023
4. Michael Hjorth, Albatros Expeditions—Permit No. 2022–024

Erika N. Davis,
Program Specialist, Office of Polar Programs.
[FR Doc. 2021–26671 Filed 12–8–21; 8:45 am]
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NATIONAL SCIENCE FOUNDATION

Notice of Permits Issued Under the Antarctic Conservation Act of 1978

AGENCY: National Science Foundation.
ACTION: Notice of permit issued.

SUMMARY: The National Science Foundation (NSF) is required to publish notice of permits issued under the Antarctic Conservation Act of 1978. This is the required notice.

FOR FURTHER INFORMATION CONTACT: Polly Penhale, ACA Permit Officer, Office of Polar Programs, National Science Foundation, 2415 Eisenhower Avenue, Alexandria, VA 22314; 703–292–8030; email: ACApermits@nsf.gov.