information in support of the claim to Laura Bryant, Gilcrease Museum, 1400 N Gilcrease Museum Road, Tulsa, OK 74127, telephone (918) 596–2747, email *laura-bryant@utulsa.edu,* by September 24, 2021. After that date, if no additional claimants have come forward, transfer of control of the sacred objects and/or objects of cultural patrimony to the Minnesota Chippewa Tribe, Minnesota (Mille Lacs Band) may proceed.

The Gilcrease Museum is responsible for notifying the Minnesota Chippewa Tribe, Minnesota (Mille Lacs Band) that this notice has been published.

Dated: August 11, 2021.

Melanie O'Brien,

Manager, National NAGPRA Program.

[FR Doc. 2021–18270 Filed 8–24–21; 8:45 am]

BILLING CODE 4312–52–P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 337–TA–1167]

Certain Laparoscopic Surgical Staplers, Reload Cartridges, and Components Thereof; Commission Determination To Review in Part a Final Initial Determination Finding a Violation of Section 337; Schedule for Filing Written Submissions on the Issues Under Review and on Remedy, Public Interest, and Bonding

AGENCY: U.S. International Trade Commission.

ACTION: Correction of notice.

SUMMARY: Correction is made to notice 86 FR 46882, which was published on August 20, 2021. There is a typographical error in the investigation number on the first page caption section. The correct investigation number should read: Investigation No. 337–TA–1167.

By order of the Commission.

Issued: August 20, 2021.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2021–18313 Filed 8–24–21; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the National Cooperative Research and Production Act of 1993—Dynamic Spectrum Alliance, Inc.

Notice is hereby given that, on August 10, 2021, pursuant to Section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (''the Act''), Dynamic Spectrum Alliance, Inc. has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing changes in its membership. The notifications were filed for the purpose of extending the Act's provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances. Specifically, Apple, Cupertino, CA, and Strathmore University, Nairobi, KENYA have been added as parties to this venture.

No other changes have been made in either the membership or planned activity of the group research project. Membership in this group research project remains open, and Dynamic Spectrum Alliance, Inc. intends to file additional written notifications disclosing all changes in membership.

On September 1, 2020, Dynamic Spectrum Alliance, Inc. filed its original notification pursuant to Section 6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to Section 6(b) of the Act on September 18, 2020 (85 FR 58390).

The last notification was filed with the Department on May 4, 2021. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on May 25, 2021 (86 FR 28150).

Suzanne Morris,

Chief, Premerger and Division Statistics, Antitrust Division.

[FR Doc. 2021–18264 Filed 8–24–21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–872]

Bulk Manufacturer of Controlled Substances Application: Benuvia Therapeutics Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Benuvia Therapeutics Inc., 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 file written comments on or objections to the issuance of the proposed registration on or before October 25, 2021. Such persons may also file a written request for a hearing on the application on or before October 25, 2021.

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

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on July 1, 2021, Benuvia Therapeutics Inc., 2700 Oakmont Drive, Round Rock, Texas 78665, 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

The company plans to bulk manufacture the listed controlled substances for the internal use intermediates or for sale to its customers. The company plans to manufacture the above-listed controlled substances in bulk to produce finished dosage forms and conduct research to develop new drug products and for clinical studies. In reference to drug codes 7360 (Marihuana), and 7370 (Tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetic. No other activities for these drug codes are authorized for this registration.

Brian S. Besser,

Acting Assistant Administrator.

[FR Doc. 2021–18234 Filed 8–24–21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–864]

Importer of Controlled Substances Application: Cedarburg Pharmaceuticals

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Cedarburg Pharmaceuticals 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 September 24, 2021. Such persons may also file a written request for a hearing on the application on or before September 24, 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 January 20, 2021, Cedarburg Pharmaceuticals, 870 Badger Circle Drive, Grafton, Wisconsin 53024–9436, 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
Marihuana Extract	7350	I
Marihuana	7360	I

The company plans to import Sodium Oxybate (derivative of Gamma-Hydroxybutyric Acid) to support Euticals Inc. post procurement quota grand. The cannabidiol from Marihuana and Marihuana Extracts is intended for analytical purposes with tetramethylpyrazine. 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. 2021–18233 Filed 8–24–21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–890]

Importer of Controlled Substances Application: Johnson Matthey Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Johnson Matthey 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 September 24, 2021. Such persons may also file a written request for a hearing on the application on or before September 24, 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 July 23, 2021, Johnson Matthey Inc., 2003 Nolte Drive, West Deptford, New Jersey 08066–1742, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Nabilone	7379	II

The company plans to import Nabilone (7379) in order to accept the return of this controlled substance from a foreign customer who no longer has a demand for this substance. No other activity for this drug code is authorized for this registration.

Brian S. Besser,

Acting Assistant Administrator.

[FR Doc. 2021–18238 Filed 8–24–21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–860]

Bulk Manufacturer of Controlled Substances Application: Absolute Standards, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Absolute Standards, Inc., 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 file written comments on or objections to the issuance of the proposed registration on or before October 25, 2021. Such persons may also file a written request for a hearing on the application on or before October 25, 2021.

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

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on May 31, 2021, Absolute Standards, Inc., 44 Rossotto Drive, Hamden, Connecticut 06514–1335, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance:

Controlled substance	Drug code	Schedule
Pentobarbital	2270	II

The company plans to bulk manufacture the listed controlled substances for internal use and for sale to its customers. No other activities for these drug codes are authorized for this registration.

Brian S. Besser,

Acting Assistant Administrator.

[FR Doc. 2021–18230 Filed 8–24–21; 8:45 am]

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