Management Ltd, London, UNITED KINGDOM; Ameren Services, St. Louis, MO; Anterix, Woodland Park, NJ; ATT Business, Dallas, TX; BEC Technologies, Richardson, TX; Blinq Network, Markham, CANADA; Bridgewater Group Consulting, Irvine, CA; Burns and McDonnell, Kansas City, MO; Cisco, San Jose, CA; Council Rock, Rochester, NY; CrescoNet, San Francisco, CA; Encore Networks, Chantilly, VA; Ericsson, Plano, TX; Evergy, Kansas City, MO; GE Industrial Communications, Rochester, NY; Hitachi-ABB, Raleigh, NC; JEA, Jacksonville, FL; K and A Engineering, White Plains, NY; L3 Harris, Rochester, NY; Mimomax Wireless, Christchurch, NEW ZEALAND; Motorola Solutions, Chicago, IL; Multi Tech Systems, Mounds View, MN; Nokia, Coppell, TX; NovaTech, Lenexa, KS; National Rural Telecommunications Cooperative, Herndon, VA; NY Power Authority (NYPA), White Plains, NY; Palmetto Technology Associates, Kiawah Island, SC; Puloli, Inc., San Francisco, CA; Southern California Edison, Rosemead, CA; SouthernLinc, Atlanta, GA; Telit Wireless Solutions, Durham, NC; World Wide Technology, Maryland Heights, MO; and Xcel Energy, Minneapolis, MN.

UBBA was formed as a Delaware non-stock member corporation. The general area of UBBA's planned activity is to empower utilities and ecosystem partners to champion the development of utility broad-band networks as a key enabler of the utility of the future, and to undertake such other activities as may from time to time be appropriate to further the purposes and achieve the goals set forth above.

Membership in UBBA remains open and UBBA intends to file additional written notifications disclosing all changes in membership.

Suzanne Morris,

Chief, Premerger and Division Statistics, Antitrust Division.

[FR Doc. 2021–12180 Filed 6–9–21; 8:45 am]

BILLING CODE 4410–11–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–847]

Importer of Controlled Substances Application: Cambrex Charles City

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Cambrex Charles City 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 July 12, 2021. Such persons may also file a written request for a hearing on the application on or before July 12, 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 request 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 May 6, 2021, Cambrex Charles City, 1205 11th Street, Charles City, Iowa 50616–3466, 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-Piperidine (ANPP).	8333	II
Phenylacetone	8501	II
Coca leaves	9040	II
Opium, raw	9600	II
Poppy Straw Concentrate.	9670	II

The company plans to import the listed controlled substances for internal use and to bulk manufacture other controlled substances in Active Pharmaceutical Ingredient (API) form for distribution to its customers. No other activity for these drug codes is authorized for this registration.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2021–12215 Filed 6–9–21; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–844]

Importer of Controlled Substances Application: Fisher Clinical Services, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Fisher Clinical Services, Inc. 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 July 12, 2021. Such persons may also file a written request for a hearing on the application on or before July 12, 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 request 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 May 17, 2021, Fisher Clinical Services, Inc. 7554 Schantz Road, Allentown, Pennsylvania 18106–9032 applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Marihuana Extract	7350	I

The company plans to import the listed controlled substances for clinical trails only. 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 the Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2021–12212 Filed 6–9–21; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–845]

Bulk Manufacturer of Controlled Substances Application: Bulk Manufacturer of Marihuana: Maridose, 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 August 9, 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–845 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). In addition to seeking to produce marihuana extract, this applicant is separately seeking to cultivate marihuana. See Notice of Application, Bulk Manufacturers of Marihuana, 84 FR 44920, 44922 (Aug. 27, 2019). DEA thus 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 May 6, 2021, Maridose, LLC., 74 Orion Street, Brunswick, Maine 04011, 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

William T. McDermott,

Assistant Administrator.

[FR Doc. 2021–12213 Filed 6–9–21; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–843]

Importer of Controlled Substances Application: National Center for Natural Products Research

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: National Center for Natural Products Research 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 July 12, 2021. Such persons may also file a written request for a hearing on the application on or before July 12, 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 request 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 April 14, 2021, National Center for Natural Products Research, 806 Hathorn Road, 135 Coy Waller Lab, University, Mississippi 38677–1848, applied to be registered as an importer 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 acquire new genetic materials with improved Cannabinoids for research and manufacturing purposes. 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 FDA-approved or non-