

Procedure, 19 CFR 210.10(b)(1), the plain language description of the accused products or category of accused products, which defines the scope of the investigation, is “powered glider-recliner mechanisms; powered rocker-recliner mechanisms; and finished seating units incorporating those mechanisms”;

(3) For the purpose of the investigation so instituted, the following are hereby named as parties upon which this notice of investigation shall be served:

- (a) The complainants are:
Ultra-Mek, Incorporated, 487 Bombay Road, Denton, North Carolina 27239
Leggett & Platt, Incorporated, 1 Leggett Rd, Carthage, MO 64836
L&P Property Management Company, 1 Leggett Rd, Carthage, MO 64836

(b) The respondents are the following entities alleged to be in violation of section 337, and are the parties upon which the complaint is to be served:

- Jiangsu Carya Smart Home Hardware Co., Ltd., No. 2 Jinchuan Rd., Industry Zone, Daqiao Town, Jiangdu District, Yangzhou Province, China
Living Style Group Ltd., 11/F, Lifung Tower, 888 Cheung Sha Wan Road, Kowloon, Hong Kong
True Innovations & Design (USA) LLC, 2052 Alton Pkwy, Irvine, CA 92606–4905, United States
Living Style (Singapore) Pte. Ltd., 3 Kallang Junction #05–02, Singapore, 339265, Singapore
Living Style (Vietnam) Ltd., 1st & 2nd Floor River Garden 170 Nguyen, Van Huong Ho Chi Minh, Vietnam 71300–0
Henglin Home Furnishings Co., Ltd., No. 378 and 380 Jiaxi Rd., Dipu Street, Huzhou, Zhejiang, 313300, China
Nanjing Hengning Home Furnishings Co., Ltd., Room 206, Building B, Shimao Chengpin, Yuhutai District, Nanjing, China
Zhejiang Hengjian Home Furnishing Co., Ltd., North Building 31 8 Baishi Lane, Gongshu District, Hangzhou, Zhejiang Hangzhou, China
Colamy, Inc., 10985 Oleander Avenue, Fontana, CA 92337
Aurora Maison, Inc., 6275 Joyce Drive, Arvada, CO 80403
Aerisnexus Innovations, Inc., 2220 NW 36th Street, Oklahoma City, OK 73112

(4) For the investigation so instituted, the Chief Administrative Law Judge, U.S. International Trade Commission, shall designate the presiding Administrative Law Judge.

The Office of Unfair Import Investigations will not participate as a party in this investigation.

The Commission is interested in the development of a thorough record on domestic industry in this investigation to facilitate a holistic review of all relevant considerations. Accordingly, the presiding administrative law judge may wish to consider what information will be necessary to make a determination with respect to complainant’s domestic industry allegations under subparagraphs (A), (B), and (C) of section 337(a)(3) based on an industry that exists or is in the process of being established; including the extent to which these allegations rely on expenditures made by third parties; and the extent to which expenditures are made outside the United States related to the domestic industry article(s) by any entity.

Responses to the complaint and the notice of investigation must be submitted by the named respondents in accordance with section 210.13 of the Commission’s Rules of Practice and Procedure, 19 CFR 210.13. Pursuant to 19 CFR 201.16(e) and 210.13(a), such responses will be considered by the Commission if received not later than 20 days after the date of service by the Commission of the complaint and the notice of investigation. Extensions of time for submitting responses to the complaint and the notice of investigation will not be granted unless good cause therefor is shown.

Failure of a respondent to file a timely response to each allegation in the complaint and in this notice may be deemed to constitute a waiver of the right to appear and contest the allegations of the complaint and this notice, and to authorize the administrative law judge and the Commission, without further notice to the respondent, to find the facts to be as alleged in the complaint and this notice and to enter an initial determination and a final determination containing such findings, and may result in the issuance of an exclusion order or a cease and desist order or both directed against the respondent.

By order of the Commission.
Issued: September 23, 2026.

Lisa Barton,
Secretary to the Commission.
[FR Doc. 2026–19713 Filed 9–25–26; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1777]

Bulk Manufacturer of Controlled Substances Application: Cargill, Incorporated

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Cargill, Incorporated 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 27, 2026. Such persons may also file a written request for a hearing on the application on or before November 27, 2026.

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 August 18, 2026, Cargill, Incorporated, 17540 Monroe Wapello Road, Eddyville, Iowa 52553, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid.	2010	I

The company plans to bulk manufacture butanediol as a raw material for industrial and consumer products. Gamma Hydroxybutyric Acid will be manufactured as a byproduct and an impurity waste of butanediol.

The company does not plan to bulk manufacture this drug. No other activity for this drug code is authorized for this registration.

Justin Wood,

Deputy Assistant Administrator.

[FR Doc. 2026–19795 Filed 9–25–26; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1778]

Bulk Manufacturer of Controlled Substances Application: Irvine Labs, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Irvine Labs, 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 submit electronic comments on or objections to the issuance of the proposed registration on or before November 27, 2026. Such persons may also file a written request for a hearing on the application on or before November 27, 2026.

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 August 13, 2026, Irvine Labs, Inc., 7305 Murdy Circle, Huntington Beach, California 92647–3533, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Ibogaine	7260	I
Lysergic acid diethylamide.	7315	I
Mescaline	7381	I
Peyote	7415	I
Diethyltryptamine	7434	I
Dimethyltryptamine	7435	I
Psilocybin	7437	I
Psilocyn	7438	I

The company plans to bulk manufacture the above listed controlled substances for research and development purposes internally and for distribution to its research customers. No other activities for these drug codes are authorized for this registration.

Justin Wood,

Deputy Assistant Administrator.

[FR Doc. 2026–19803 Filed 9–25–26; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1767]

Importer of Controlled Substances Application: Benuvia Operations, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Benuvia Operations, 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.

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 28, 2026. Such persons may also file a written request for a hearing on the application on or before October 28, 2026.

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

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on August 11, 2026, Benuvia Operations, LLC., 3950 North Mays Street, Round Rock, Texas 78665–2729, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Ibogaine	7260	I

The company plans to import the listed controlled substance to support internal research and development toward the bulk manufacture of API for future clinical trials. 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.

Justin Wood,

Deputy Assistant Administrator.

[FR Doc. 2026–19792 Filed 9–25–26; 8:45 am]

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NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[NASA Document Number: 26–056; NASA Docket Number: NASA–2026–0463]

Name of Information Collection: NASA Generic Clearance for Conferences, Meetings, Workshops, Registrations and Challenges

AGENCY: National Aeronautics and Space Administration (NASA).

ACTION: Notice of new information collection.