

Device (Vietnam) Company, Limited of Binh Duong Province, Vietnam; TCL Smart Screen Technology HK of New Territories, Hong Kong; TCL Moka International Ltd. of New Territories, Hong Kong; TTE Technology, Inc. of Irvine, California; Hisense Co., Ltd. of Shandong Province, China; Hisense USA Corporation of Suwanee, Georgia; and Hisense Electronics Manufacturing Company of America Corporation of Suwanee, Georgia (collectively, "Respondents"). The Office of Unfair Import Investigations is not participating in this investigation. *Id.* at 1674.

On July 10, 2026, non-party Dolby Laboratories, Inc. ("Dolby") filed a motion pursuant to Commission Rule 210.19, 19 CFR 210.19, to intervene as an intervenor with full participation rights as to the '168 and '751 patents. On July 22, 2026, Complainants filed an opposition to the motion. Dolby filed a reply in support of its motion on July 27, 2026. While no other responses were received, Dolby certifies that the Respondents do not oppose the motion.

On August 12, 2026, the ALJ issued the subject ID (Order No. 17) granting Dolby's motion. The ALJ has determined that Dolby should be granted intervenor status in this investigation, but not status as a respondent. The ID states that Dolby's participation in this investigation should begin with the issuance of the subject ID and that Dolby may participate as an intervenor, including with respect to all claims and defenses at issue in the investigation regarding the '168 and '751 patents. No petitions for review were filed.

The Commission has determined not to review the subject ID. The Commission vote for this determination took place on September 9, 2026.

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: September 9, 2026.

**Lisa Barton,**

*Secretary to the Commission.*

[FR Doc. 2026-18715 Filed 9-11-26; 8:45 am]

**BILLING CODE 7020-02-P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-1763]

#### Importer 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 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 14, 2026. Such persons may also file a written request for a hearing on the application on or before October 14, 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 13, 2026, Irvine Labs, Inc., 7305 Murdy Circle, Huntington Beach, California 92647-3533, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

| Controlled substance        | Drug code | Schedule |
|-----------------------------|-----------|----------|
| Lysergic acid diethylamide. | 7315      | I        |
| Marihuana Extract .....     | 7350      | I        |
| Marihuana .....             | 7360      | I        |
| Tetrahydrocannabinols ....  | 7370      | I        |
| Mescaline .....             | 7381      | I        |
| Peyote .....                | 7415      | I        |
| Diethyltryptamine .....     | 7434      | I        |
| Dimethyltryptamine .....    | 7435      | I        |
| Psilocybin .....            | 7437      | I        |
| Psilocyn .....              | 7438      | I        |

The company plans to import bulk substances to support internal research, clinical trials, analytical purposes, and distribution to their customers. In reference to drug codes Marihuana Extract (7350), Marihuana (7360) and Tetrahydrocannabinols (7370), the company plans to import a raw plant material and extracts. No other activities for these drug codes are 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. 2026-18757 Filed 9-11-26; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-1759]

#### Bulk Manufacturer of Controlled Substances Application: Cambrex High Point, Inc.

**AGENCY:** Drug Enforcement Administration, Justice.

**ACTION:** Notice of application.

**SUMMARY:** Cambrex High Point, 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 13, 2026. Such persons may also file a written request for a hearing on the application on or before November 13, 2026.