public hearing in connection with the review, originally scheduled for April 27, 2023, was cancelled (88 FR 26598, April 25, 2023).

The Commission made this determination pursuant to section 751(c) of the Act (19 U.S.C. 1675(c)). It completed and filed its determination in this review on June 26, 2023. The views of the Commission are contained in USITC Publication 5433 (June 2023), entitled *Dioctyl Terephthalate from South Korea: Investigation No. 731–TA–1330 (Review).*

By order of the Commission.

Issued: June 26, 2023.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2023–13862 Filed 6–28–23; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1197]

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 July 31, 2023. Such persons may also file a written request for a hearing on the application on or before July 31, 2023.

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

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on April 20, 2023, 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 the bulk substances to support internal research, clinical trials, analytical purposes, and distribution to their customers. In reference to drug codes 7360 (Marihuana), 7350 (Marihuana Extract), and 7370 (Tetrahydrocannabinols) 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.

Matthew Strait,

Deputy Assistant Administrator.

[FR Doc. 2023–13812 Filed 6–28–23; 8:45 am]

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DEPARTMENT OF LABOR

Office of Workers' Compensation Programs

[OMB Control No. 1240–0021]

Proposed Extension of Existing Collection; Comment Request

AGENCY: Office of Workers' Compensation Programs, Labor.

ACTION: Request for public comment.

SUMMARY: The Department of Labor, as part of its continuing effort to reduce paperwork and respondent burden, conducts a pre-clearance request for comment to provide the general public and Federal agencies with an opportunity to comment on proposed collections of information in accordance with the Paperwork Reduction Act of 1995. This request helps to ensure that: requested data can be provided in the desired format; reporting burden (time and financial resources) is minimized; collection instruments are clearly understood; and the impact of collection requirements on respondents can be properly assessed. Currently, OWCP is soliciting comments on the information collection for the Provider Enrollment Form (PE–1168).

DATES: All comments must be received on or before August 28, 2023.