The company plans to import analytical reference standards for distribution to its customers for research and analytics purposes. Placement of these drug codes onto the company's registration does not translate into automatic approval of subsequent permit applications to import controlled substances. Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized in 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.

Claude Redd,

Acting Deputy Assistant Administrator.

[FR Doc. 2023–21404 Filed 9–28–23; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1274]

Importer of Controlled Substances Application: Cambridge Isotope Laboratories, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Cambridge Isotope Laboratories, 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 30, 2023. Such persons may also file a written request for a hearing on the application on or before October 30, 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 August 25, 2023, Cambridge Isotope Laboratories, Inc. 50 Frontage Road, Andover, Massachusetts 01810–5413, 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
Tetrahydrocannabinols	7370	I
Morphine	9300	II

The company plans to import the listed controlled substances for preparation of analytical standards and formulations. In reference to drug codes 7370 (Tetrahydrocannabinols), the company plans to import a synthetic Tetrahydrocannabinol. 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.

Claude Redd,

Acting Deputy Assistant Administrator.

[FR Doc. 2023–21402 Filed 9–28–23; 8:45 am]

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DEPARTMENT OF LABOR

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Formaldehyde Standard

ACTION: Notice of availability; request for comments.

SUMMARY: The Department of Labor (DOL) is submitting this Occupational Safety & Health Administration (OSHA)-sponsored information collection request (ICR) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995 (PRA). Public comments on the ICR are invited.

DATES: The OMB will consider all written comments that the agency receives on or before October 30, 2023.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

Comments are invited on: (1) whether the collection of information is necessary for the proper performance of the functions of the Department, including whether the information will have practical utility; (2) the accuracy of the agency's estimates of the burden and cost of the collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility and clarity of the information collection; and (4) ways to minimize the burden of the collection of information on those who are to respond, including the use of automated collection techniques or other forms of information technology.

FOR FURTHER INFORMATION CONTACT: Nicole Bouchet by telephone at 202–693–0213, or by email at *DOL_PRA_PUBLIC@dol.gov.*

SUPPLEMENTARY INFORMATION: The Formaldehyde Standard and its collections of information are designed to provide protection for workers from the adverse health effects associated with occupational exposure to formaldehyde. The Standard requires employers to monitor worker exposure and provide notification to workers of their exposure. Employers are required to make available medical surveillance to workers. For additional substantive information about this ICR, see the related notice published in the **Federal Register** on June 28, 2023 (88 FR 41984).

This information collection is subject to the PRA. A Federal agency generally cannot conduct or sponsor a collection of information, and the public is generally not required to respond to an information collection, unless the OMB approves it and displays a currently valid OMB Control Number. In addition, notwithstanding any other provisions of

law, no person shall generally be subject to penalty for failing to comply with a collection of information that does not display a valid OMB Control Number. *See* 5 CFR 1320.5(a) and 1320.6.

DOL seeks PRA authorization for this information collection for three (3) years. OMB authorization for an ICR cannot be for more than three (3) years without renewal. The DOL notes that information collection requirements submitted to the OMB for existing ICRs receive a month-to-month extension while they undergo review.

Agency: DOL–OSHA.

Title of Collection: Formaldehyde Standard.

OMB Control Number: 1218–0145.

Affected Public: Private sector—businesses or other for-profits.

Total Estimated Number of Respondents: 86,575.

Total Estimated Number of Responses: 990,175.

Total Estimated Annual Time Burden: 263,172 hours.

Total Estimated Annual Other Costs Burden: $54,153,624.

(Authority: 44 U.S.C. 3507(a)(1)(D))

Nicole Bouchet,

Acting Departmental Clearance Officer.

[FR Doc. 2023–21417 Filed 9–28–23; 8:45 am]

BILLING CODE 4510–26–P

DEPARTMENT OF LABOR

[Agency Docket Number DOL–2023–xxxx]

Efforts by Certain Foreign Countries To Eliminate the Worst Forms of Child Labor; Identify Child Labor, Forced Labor, and Forced or Indentured Child Labor in the Production of Goods in Foreign Countries; and Share Business Practices To Reduce the Likelihood of Forced Labor or Child Labor in the Production of Goods

AGENCY: The Bureau of International Labor Affairs, United States Department of Labor.

ACTION: Notice; request for information and invitation to comment.

SUMMARY: This notice is a request for information and/or comment on three reports issued by the Bureau of International Labor Affairs (ILAB) regarding child labor and forced labor in certain foreign countries. Relevant information submitted by the public will be used by the Department of Labor (DOL) in preparing its ongoing reporting as required under Congressional mandates and a Presidential directive.

DATES: Submitters of information are requested to provide their submission to DOL's Office of Child Labor, Forced Labor, and Human Trafficking (OCFT) at the email or physical address below by December 15, 2023.

ADDRESSES:

To Submit Information: Information should be submitted directly to OCFT, Bureau of International Labor Affairs, U.S. Department of Labor. Comments, identified as Docket No. DOL–2023–xxxx, may be submitted by any of the following methods:

Federal eRulemaking Portal: The portal includes instructions for submitting comments. Parties submitting responses electronically are encouraged not to submit paper copies.

Facsimile (fax): OCFT at 202–693–4830.

Mail, Express Delivery, Hand Delivery, and Messenger Service (1 copy): Matthew Fraterman, U.S. Department of Labor, OCFT, Bureau of International Labor Affairs, 200 Constitution Avenue NW, Room S–5315, Washington, DC 20210.

Email: Email submissions should be addressed to Matthew Fraterman (*Fraterman.matthew@dol.gov*).

508 Compliance: Pursuant to section 508 of the Rehabilitation Act of 1973 (29 U.S.C. 794d), as amended. Section 508 became enforceable on June 21, 2001, and the Revised 508 standards issued by the United States Access Board (36 CFR part 1194), January 2018 require that Information and Communication Technology (ICT) procured, developed, maintained, and used by Federal departments and agencies is accessible to and usable by Federal employees and members of the public including people with disabilities. All documents received in electronic format must be accessible using assistive technologies such as a screen reader, *e.g.,* Job Aid with Speech (JAWS), NonVisual Desktop Access (NVDA), ZoomText, to name a few. The product should also be navigable using other means such as a keyboard or voice commands. Accessible document formats are either Microsoft Word or equivalent and Portable Document Format with OCR.

The Department of Labor requests that your submissions through the portal comply with our DOL Policies as well as the 508 Standards as referenced above.

FOR FURTHER INFORMATION CONTACT: Matthew Fraterman (*Fraterman.matthew@dol.gov*). Telephone: 202–693–4770.

SUPPLEMENTARY INFORMATION: The 2022 Findings on the Worst Forms of Child Labor (TDA Report), published on September 26, 2023, assesses efforts of 131 countries to eliminate the worst forms of child labor in 2022 and assesses whether countries made significant, moderate, minimal, or no advancement during that year. It also suggests actions foreign countries can take to eliminate the worst forms of child labor through legislation, enforcement, coordination, policies, and social programs. The 2022 edition of the List of Goods Produced by Child Labor or Forced Labor (TVPRA List), published on September 21, 2022, makes available to the public a list of goods from countries that ILAB has reason to believe are produced by child labor or forced labor in violation of international standards, including, to the extent practicable, goods that are produced with inputs that are produced with forced labor or child labor. DOL welcomes new information on any of the goods identified on the TVPRA List. Finally, the List of Products Produced by Forced or Indentured Child Labor (E.O. List), provides a list of products, identified by country of origin, that DOL, in consultation and cooperation with the Departments of State (DOS) and Homeland Security (DHS), has a reasonable basis to believe might have been mined, produced, or manufactured with forced or indentured child labor. Relevant information submitted by the public will be used by DOL in preparing the next edition of the TDA Report, to be published in 2024; the next edition of the TVPRA List, which will also be published in 2024; and for possible updates to the E.O. List as needed.

This notice is also a request for information and/or comment on Comply Chain: Business Tools for Labor Compliance in Global Supply Chains (Comply Chain). ILAB is seeking information on current practices of firms, business associations, and other private sector groups to reduce the likelihood of child labor and forced labor in the production of goods. This information and/or comment is sought to fulfill ILAB's mandate under the Trafficking Victims Protection Reauthorization Act of 2005 (TVPRA) to work with persons who are involved in the production of goods made with forced labor or child labor. Comply Chain seeks to address this mandate through the creation of a standard set of practices for worker-driven social compliance that will reduce the likelihood that such persons will produce goods using forced labor or child labor. Comply Chain also achieves a much broader purpose by actively supporting the efforts of companies that seek to address these issues within their own supply chains. Relevant information and/or comment submitted to ILAB will be used to improve and