

Correction

In the **Federal Register** of May 17, 2022, in FR #2022–10522, on page 29870, in the second column, correct the **DATES** caption to read:

DATES: *Comments Due Date:* July 18, 2022.

Nacheshia Foxx,

Federal Register Liaison for HUD, Office of Regulations, Office of the General Counsel.

[FR Doc. 2022–10946 Filed 5–20–22; 8:45 am]

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DEPARTMENT OF JUSTICE**Drug Enforcement Administration**

[Docket No. DEA–963]

Bulk Manufacturer of Controlled Substances Application: Scottsdale Research Institute

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Scottsdale Research Institute, has applied to be registered as a bulk manufacturer 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 submit electronic comments on or objections to the issuance of the proposed registration on or before July 22, 2022. Such persons may also file a written request for a hearing on the application on or before July 22, 2022.

ADDRESSES: The DEA 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 January 5, 2022, 5436 East Tapekim Road, Cave Creek, Arizona 85331, applied to be registered as a bulk manufacturer 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 bulk manufacture the listed controlled substances for the internal research use and for distribution to customers for research purposes. No other activities for these drug codes are authorized for this registration.

Matthew J. Strait,

Deputy Assistant Administrator.

[FR Doc. 2022–10973 Filed 5–20–22; 8:45 am]

BILLING CODE P

DEPARTMENT OF LABOR

Agency Information Collection Activities; Submission for OMB Review; Comment Request; H–2B Application for Temporary Employment Certification

ACTION: Notice of availability; request for comments.

SUMMARY: The Department of Labor (DOL) is submitting this Employment and Training Administration (ETA)-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 June 22, 2022.

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) if the information will be processed and used in a timely manner; (3) 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; (4)

ways to enhance the quality, utility and clarity of the information collection; and (5) 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: Mara Blumenthal by telephone at 202–693–8538, or by email at DOL_PRA_PUBLIC@dol.gov.

SUPPLEMENTARY INFORMATION: This information collection is required by Sections 101(a)(15)(H)(ii)(b) and 214(c) of the Immigration and Nationality Act (INA) (8 U.S.C. 1011(a)(15)(H)(ii)(b) and 1184(c)), as well as 8 CFR 214.2(h)(6), 20 CFR part 655, subpart A, and 29 CFR part 503. This ICR supports Department of Labor and Department of Homeland Security regulations that contain information collections under the H–2B labor certification program. The H–2B program enables employers to bring nonimmigrant foreign workers to the United States to perform non-agricultural work of a temporary nature. The information contained in the Form ETA–9142B, H–2B Application for Temporary Employment Certification, and corresponding appendices serve as the basis for the Secretary’s determination that qualified U.S. workers are not available to perform the services or labor needed by the employer and that the wages and working conditions of similarly employed U.S. workers will not be adversely affected by the employment of H–2B workers. For additional substantive information about this ICR, see the related notice published in the **Federal Register** on January 12, 2022 (87 FR 1787).

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–ETA.