1021, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Cocaine	9041	II
Ecgonine	9180	II

The company plans to bulk manufacture the listed controlled substances for use as internal intermediates or for sale to its customers. No other activities for these drug codes are authorized for this registration.

Matthew Strait,

Deputy Assistant Administrator.

[FR Doc. 2025–06706 Filed 4–17–25; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1531]

Bulk Manufacturer of Controlled Substances Research Triangle Institute

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Research Triangle Institute 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 June 17, 2025. Such persons may also file a written request for a hearing on the application on or before June 17, 2025.

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 March 14, 2025, Research Triangle Institute, 3040 East Cornwallis Road, Hermann Building Room 106, Research Triangle Park, North Carolina 27709–2194, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Tetrahydrocannabinols	7370	I

The company plans to bulk manufacture the listed controlled substance(s) synthetically for distribution to its customers for research and as analytical reference standards. No other activity for this drug code is authorized for this registration.

Matthew Strait,

Deputy Assistant Administrator.

[FR Doc. 2025–06710 Filed 4–17–25; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1530]

Bulk Manufacturer of Controlled Substances Application: Restek Corporation

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Restek Corporation 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 June 17, 2025. Such persons may also file a written request for a hearing on the application on or before June 17, 2025.

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 February 19, 2025, Restek Corporation, 110 Benner Circle, Bellefonte, Pennsylvania 16823–8433, 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
Methaqualone	2565	I
JWH–018 (also known as AM678) (1-Pentyl-3-(1naphthoyl)indole)	7118	I
JWH–200 (1-[2-(4-Morpholinyl)ethyl]-3-(1- naphthoyl)indole	7200	I
CP–47,497 (5-(1,1-Dimethylheptyl)-2-[(1R,3S)-3-hydroxycyclohexy]-phenol)	7297	I
CP–47,497 C8 Homologue (5-(1,1-Dimethyloctyl)-2- [(1R,3S)3-hydroxycyclohexyl]-phenol)	7298	I
Lysergic acid diethylamide	7315	I
Marihuana	7360	I
Tetrahydrocannabinols	7370	I
4-Methyl-2,5-dimethoxyamphetamine	7395	I
3,4-Methylenedioxyamphetamine	7400	I

Controlled substance	Drug code	Schedule
3,4-Methylenedioxy-N-ethylamphetamine	7404	I
3,4-Methylenedioxymethamphetamine	7405	I
Bufotenine	7433	I
Psilocybin	7437	I
Psilocyn	7438	I
Cyprenorphine	9054	I
Dihydromorphine	9145	I
Heroin	9200	I
Normorphine	9313	I
Beta-hydroxyfentanyl	9830	I
Beta-hydroxy-3-methylfentanyl	9831	I

The company plans to bulk manufacture the listed controlled substances for the Drug Enforcement Administration-exempted certified reference materials. In-house synthesis gives access to compounds that are difficult to source. In reference to drug codes 7360 (Marihuana), and 7370 (Tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetic. No other activities for these drug codes are authorized for this registration.

Matthew Strait,

Deputy Assistant Administrator.

[FR Doc. 2025–06709 Filed 4–17–25; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Notice of Availability; Service Contract Inventory

AGENCY: Justice Management Division, Department of Justice.

ACTION: Notice of availability.

SUMMARY: The Justice Management Division (JMD), Department of Justice (DOJ) is publishing this notice to advise the public of the availability of its FY 2023 Service Contracts Inventory and Inventory Supplement.

ADDRESSES: *https://www.justice.gov/jmd/service-contract-inventory*.

FOR FURTHER INFORMATION CONTACT: Wendy G. Devoe, Office of Acquisition Management, Justice Management Division, U.S. Department of Justice, Washington, DC 20530; Phone: 202–285–7425; Email: *Wendy.Devoe@usdoj.gov*.

SUPPLEMENTARY INFORMATION: The inventory includes service contract actions over $25,000 that were awarded in Fiscal Year (FY) 2023. Additionally, the inventory supplement includes information collected from contractors on the amount invoiced and direct labor hours expended for covered service contracts. The Department of Justice analyzes this data for the purpose of determining whether its contract labor is being used in an effective and appropriate manner and if the mix of federal employees and contractors in the agency is effectively balanced. The inventory and supplement do not include contractor proprietary or sensitive information.

Authority: Section 743 of Division C of the FY 2010 Consolidated Appropriations Act, Pub. L. 111–117.

Dated: April 15, 2025.

Darwin Arceo,

Department Clearance Officer for PRA, U.S. Department of Justice.

[FR Doc. 2025–06708 Filed 4–17–25; 8:45 am]

BILLING CODE 4410–02–P

DEPARTMENT OF LABOR

Agency Information Collection Activities; Submission for OMB Review; Comment Request; National Safety Stand-Down To Prevent Falls in Construction

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 May 19, 2025.

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.

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

SUPPLEMENTARY INFORMATION: The National Safety Stand-Down to Prevent Falls in construction raises fall hazard awareness across the country in an effort to stop fall fatalities and injuries. Participants (mainly employers) download a Certificate of Participation by completing a simple eight question online survey. The survey is the primary means that OSHA will have for validating participation in the Stand-Down. For additional substantive information about this ICR, see the related notice published in the **Federal Register** on November 5, 2024 (89 FR 87897).

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.

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