

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.*

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**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.*

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**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-(1-naphthoyl)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-hydroxycyclohexyl]-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