Controlled substance	Drug code	Schedule
Nabilone	. 7379	II
1-Phenylcyclohexylamine	. 7460	II
Phencyclidine	. 7471	II
ANPP (4-Anilino-N-phenethyl-4-piperidine)		П
Phenylacetone		П
1-Piperidinocyclohexanecarbonitrile		П
Alphaprodine		П
Anileridine		П
Cocaine		lii
Codeine		ii
Etorphine HCI		ii
Dihydrocodeine		l ii
Oxycodone		lii
Hydromorphone		lii
		l ii
Diphenoxylate		li Ii
Ecgonine		
Ethylmorphine		II
Hydrocodone		II.
Levomethorphan		ii ii
Levorphanol		II
Isomethadone		II
Meperidine		II
Meperidine-intermediate-A		II
Meperidine intermediate-B	. 9233	II
Meperidine intermediate-C	. 9234	II
Metazocine	. 9240	II
Methadone	. 9250	II
Methadone intermediate	. 9254	II
Metopon	. 9260	П
Dextropropoxyphene, bulk (non-dosage forms)		П
Morphine		П
Oripavine		П
Thebaine		l ii
Dihydroetorphine		ii
Levo-alphacetylmethadol		l ii
Oxymorphone		l ii
Noroxymorphone		ii
Phenazocine		ii
		l ii
Thiafentanil		li Ii
Piminodine		
Racemethorphan		II
Racemorphan		II
Alfentanil		II
Remifentanil		II
Sufentanii		II
Carfentanil		II.
Tapentadol		II
Bezitramide		II
Fentanyl		II
Moramide-intermediate	. 9802	II

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. 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.

Marsha Ikner,

Acting Deputy Assistant Administrator. [FR Doc. 2024–09802 Filed 5–3–24; 8:45 am] BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1349]

Bulk Manufacturer of Controlled Substances Application: 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 July 5, 2024. Such persons may also file a written request for a hearing on the application on or before July 5, 2024.

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 11, 2024, 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	1

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

Marsha Ikner,

Acting Deputy Assistant Administrator. [FR Doc. 2024–09783 Filed 5–3–24; 8:45 am] BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1359]

Importer of Controlled Substances Application: Restek Corporation

AGENCY: Drug Enforcement Administration, Justice. **ACTION:** Notice of application.

SUMMARY: Restek Corporation 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 June 5, 2024. Such persons may also file a written request for a

hearing on the application on or before June 5, 2024.

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 March 5, 2024, Restek Corporation, 110 Benner Circle, Bellefonte, Pennsylvania 16823–8433, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Amineptine	1219	ı
Amineptine Mesocarb	1227	1
3-Fluoro-N-methylcathinone (3–FMC)	1233	1
Cathinone	1235	1
Methcathinone	1237	1
4-Fluoro-N-methylcathinone (4–FMC)	1238	1
Para-Methoxymethamphetamine (PMMA), 1-(4-1245 I N methoxyphenyl)-N-methylpropan-2-amine	1245	1
Pentedrone (a-methylaminovalerophenone)	1246	1
Mephedrone (4-Methyl-N-methylcathinone	1248	1
4-Methyl-N-ethylcathinone (4-MEC)	1249	1
	1258	1
Naphyrone	1475	1
Methiopropamine (N-methyl-1-(thiophen-2-yl)propan-2- 1478 I N amine)	1478	1
N,N-Dimethylamphetamine	1480	1
Fenethylline	1503	1
Aminorex	1585	1
4-Methylaminorex (cis isomer)	1590	1
4,4'-Dimethylaminorex	1595	1
Gamma Hydroxybutyric Acid	2010	1
Methaqualone	2565	1
Mecloqualone	2572	1
JWH-250 (1-Pentyl-3-(2-methoxyphenylacetyl) indole)	6250	1
SR-18 (Also known as RCS-8) (1-Cyclohexylethyl-3-(2-methoxyphenylacetyl) indole)	7008	1
ADB-FÙBINACA (N-(1-amino-3,3-dimethyl-1-oxobutan-2-yl)-1-(4-fluorobenzyl)-1H-indazole-3-carboxamide)	7010	1
5-Fluoro-UR-144 and XLR11 [1-(5-Fluoro-pentyl)1H-indol-3-yl](2,2,3,3-tetramethylcyclopropyl)methanone		1