

Controlled substance	Drug code	Schedule
Lysergic acid diethylamide .....	7315	I
Marihuana Extract .....	7350	I
Marihuana .....	7360	I
Tetrahydrocannabinols .....	7370	I
Mescaline .....	7381	I
JWH-398 (1-Pentyl-3-(4-chloro-1-naphthoyl) indole) .....	7398	I
3,4-Methylenedioxymethamphetamine .....	7405	I
5-Methoxy-N-N-dimethyltryptamine .....	7431	I
Psilocyn .....	7438	I
4-Methyl-alpha-pyrrolidinopropiophenone (4-MePPP) .....	7498	I
MDPV (3,4-Methylenedioxypropylvalerone) .....	7535	I
Methylone (3,4-Methylenedioxy-N-methylcathinone) .....	7540	I
Butylone .....	7541	I
Pentylone .....	7542	I
Codeine-N-oxide .....	9053	I
Desomorphine .....	9055	I
Dihydromorphine .....	9145	I
Heroin .....	9200	I
Morphine-N-oxide .....	9307	I
Normorphine .....	9313	I
Tilidine .....	9750	I
Alpha-methylfentanyl .....	9814	I
Acetyl Fentanyl (N-(1-phenethylpiperidin-4-yl)-N-phenylacetamide) .....	9821	I
Fentanyl-related substance .....	9850	I
Methamphetamine .....	1105	II
Phenmetrazine .....	1631	II
Methylphenidate .....	1724	II
Amobarbital .....	2125	II
Pentobarbital .....	2270	II
Secobarbital .....	2315	II
Glutethimide .....	2550	II
Phencyclidine .....	7471	II
4-Anilino-N-phenethyl-4-piperidine (ANPP) .....	8333	II
Norfentanyl .....	8366	II
Phenylacetone .....	8501	II
Codeine .....	9050	II
Dihydrocodeine .....	9120	II
Oxycodone .....	9143	II
Hydromorphone .....	9150	II
Diphenoxylate .....	9170	II
Ecgonine .....	9180	II
Ethylmorphine .....	9190	II
Hydrocodone .....	9193	II
Levorphanol .....	9220	II
Meperidine .....	9230	II
Meperidine intermediate-A .....	9232	II
Meperidine intermediate-B .....	9233	II
Meperidine intermediate-C .....	9234	II
Methadone intermediate .....	9254	II
Dextropropoxyphene, bulk (non-dosage forms) .....	9273	II
Morphine .....	9300	II
Thebaine .....	9333	II
14-Hydroxymorphine .....	9665	II
Noroxymorphone .....	9668	II
Sufentanil .....	9740	II
Fentanyl .....	9801	II

The company plans to import the listed controlled substances for sale to research facilities for drug testing and analysis. In reference to drug codes 7360 (Marihuana) and 7370 (Tetrahydrocannabinols) the company plans to import a synthetic cannabidiol and 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.

**Matthew J. Strait,**

*Deputy Assistant Administrator.*

[FR Doc. 2022-09057 Filed 4-27-22; 8:45 am]

**BILLING CODE 4410-09-P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-992]

### Importer of Controlled Substances Application: Purisys, LLC

**AGENCY:** Drug Enforcement Administration, Justice.

**ACTION:** Notice of application.

**SUMMARY:** Purisys, LLC has applied to be registered as an importer 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 May 31, 2022. Such persons may also file a written request for a hearing on the application on or before May 31, 2022.

**ADDRESSES:** The Drug Enforcement Administration (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. All requests for a hearing must be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on January 21, 2022, Purisys, LLC, 1550 Olympic Drive, Athens, Georgia 30601-1602, applied to be registered as an importer 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
Nabilone .....	7379	II
Phenylacetone .....	8501	II
Levorphanol .....	9220	II
Thebaine .....	9333	II
Opium, Raw .....	9600	II
Opium, Power .....	9639	II
Opium Granulated .....	9640	II
Noroxymorphone .....	9668	II

Controlled substance	Drug code	Schedule
Concentrate of Poppy Straw.	9670	II
Tapentadol .....	9780	II

The company plans to import Opium, Raw (9600), Opium, Powered (9639) and Opium, Granulated (9640) to manufacture Active Pharmaceutical Ingredient (API) only for distribution to its customers. The company plans to import Phenylacetone (8501) and Poppy Straw Concentrate (9670), to bulk manufacture other controlled substances for distribution to its customers. The company plans to import impurities of buprenorphine that have been determined by DEA to be captured under Thebaine (9333). In reference to Marihuana (7360) and Tetrahydrocannabinols (7370) the company plans to import as synthetic. No other activity for these drug codes is 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.

**Matthew J. Strait,**

*Deputy Assistant Administrator.*

[FR Doc. 2022-09034 Filed 4-27-22; 8:45 am]

**BILLING CODE 4410-09-P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-993]

#### Importer of Controlled Substances Application: SpecGX LLC

**AGENCY:** Drug Enforcement Administration, Justice.

**ACTION:** Notice of application.

**SUMMARY:** SpecGX, LLC has applied to be registered as an importer 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 May 31, 2022. Such persons may also file a written request for a hearing on the application on or before May 31, 2022.

**ADDRESSES:** The Drug Enforcement Administration (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. All requests for a hearing must be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on January 26, 2022, SpecGX LLC, 3600 North 2nd Street, Saint Louis, Missouri 63147, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Marihuana .....	7360	I
Phenylacetone .....	8501	II
Coca Leaves .....	9040	II
Thebaine .....	9333	II
Opium, Raw .....	9600	II
Poppy Straw Concentrate.	9670	II
Tapentadol .....	9780	II

The company plans to import the listed controlled substances for bulk manufacture into Active Pharmaceutical Ingredients (API) for distribution to its customers. In reference to Tapentadol (9780) and Thebaine (9333), the company plans to import intermediate forms of these controlled substances for further manufacturing prior to distribution to its customers. In reference to drug code 7360 (Marihuana), the company plans to import synthetic cannabinol. No other activity for this drug is authorized for this registration. Placement of these codes onto the company's registration