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-alphapyrrolidinopropiophenone (4-MePPP)	7498	I
MDPV (3,4-Methylenedioxypyrovalerone)	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-Hydroxmorphone	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 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 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
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 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 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