

| 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.  
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**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.  
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**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