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 January 26, 2022, SpecGx LLC, 3600 North Second Street, Saint Louis, Missouri 63147–3457, 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 |
| Tetrahydrocannabinols | 7370 | I |
| Codeine-N-oxide | 9053 | I |
| Noroxymorphone | 9145 | I |
| Difenoxin | 9168 | I |
| Morphine-N-oxide | 9307 | I |
| Normorphine | 9313 | I |
| Alphamethadol | 9605 | I |
| Betamethadol | 9609 | I |
| Norlevorphanol | 9634 | I |
| Acetyl Fentanyl (N-(1-phenethylpiperidin-4-yl)-N-phenylacetamide) | 9821 | I |
| Butyryl Fentanyl | 9822 | I |
| Fentanyl related-compounds as defined in 21 CFR 1308.11(h) | 9850 | I |
| Amphetamine | 1100 | II |
| Methamphetamine | 1105 | II |
| Lisdexamfetamine | 1205 | II |
| Methylphenidate | 1724 | II |
| Nabilone | 7379 | II |
| ANPP (4-Anilino-N-phenethyl-4-piperidine) | 8333 | II |
| Phenylacetone | 8501 | II |
| Codeine | 9050 | II |
| Dihydrocodeine | 9120 | II |
| Oxycodone | 9143 | II |
| Hydromorphone | 9150 | II |
| Diphenoxylate | 9170 | II |
| Ecgonine | 9180 | 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 | 9250 | II |
| Methadone intermediate | 9254 | II |
| Dextropropoxyphene, bulk (non-dosage forms) | 9273 | II |
| Morphine | 9300 | II |
| Oripavine | 9330 | II |
| Thebaine | 9333 | II |
| Opium tincture | 9630 | II |
| Opium, powdered | 9639 | II |
| Oxymorphone | 9652 | II |
| Noroxymorphone | 9668 | II |
| Alfentanil | 9737 | II |
| Remifentanil | 9739 | II |
| Sufentanil | 9740 | II |
| Tapentadol | 9780 | II |
| Fentanyl | 9801 | II |

The company plans to bulk manufacture the listed controlled substances for the internal use intermediates or for sale to its customers. In reference to dug code 7370 (Tetrahydrocannabinols), the company plans to bulk manufacture this drug as synthetic. No other activities for these drug codes are authorized for this registration.

**Matthew J. Strait,**

*Deputy Assistant Administrator.*

[FR Doc. 2022–09035 Filed 4–27–22; 8:45 am]

**BILLING CODE P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

**[Docket No. DEA–999]**

### Importer of Controlled Substances Application: Royal Emerald Pharmaceuticals Research and Development

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

**ACTION:** Notice of application.

**SUMMARY:** Royal Emerald Pharmaceuticals 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 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 25, 2022, Royal Pharmaceuticals Research and Development, 14011 Palm Drive, Desert Hot Springs, California 92240, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

| | | |
|---|---|---|
| Marihuana Extract .. | 7350 | I |
| Marihuana ..... | 7360 | I |
| Tetrahydrocannabinols. | 7370 | I |

The company plans to import Marihuana seeds and immature Marihuana plants in the form of Active Pharmaceutical Ingredients (API) and botanical raw materials for DEA-approved legitimate scientific medical research and/or industrial purposes.

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. 2022–09061 Filed 4–27–22; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

**[Docket No. DEA–996]**

### Importer of Controlled Substances Application: VHG Labs DBA LGC Standards

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

**ACTION:** Notice of application.

**SUMMARY:** VHG Labs DBA LGC Standards 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 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 February 18, 2022, VHG Labs DBA LGC Standards, 3 Perimeter Road, Manchester, New Hampshire 03103, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

| Controlled substance | Drug code | Schedule |
|---|---|---|
| Cathinone | 1235 | I |
| Methcathinone | 1237 | I |
| Naphyrone | 1258 | I |
| N-Ethylamphetamine | 1475 | I |
| JWH-250 (1-Pentyl-3-(2-methoxyphenylacetyl) indole) | 6250 | I |
| SR-18 (Also known as RCS-8) (1-Cyclohexylethyl-3-(2-methoxyphenylacetyl) indole) | 7008 | I |
| APINACA and AKB48 N-(1-Adamantyl)-1-pentyl-1H-indazole-3-carboxamide | 7048 | I |
| JWH-081 (1-Pentyl-3-(1-(4-methoxynaphthoyl) indole) | 7081 | I |
| SR-19 (Also known as RCS-4) (1-Pentyl-3-[(4-methoxy)-benzoyl] indole | 7104 | I |
| JWH-018 (also known as AM678) (1-Pentyl-3-(1-naphthoyl)indole) | 7118 | I |
| JWH-122 (1-Pentyl-3-(4-methyl-1-naphthoyl) indole) | 7122 | I |
| UR-144 (1-Pentyl-1H-indol-3-yl)(2,2,3,3-tetramethylcyclopropyl)methanone | 7144 | I |
| JWH-203 (1-Pentyl-3-(2-chlorophenylacetyl) indole) | 7203 | I |
| Ibogaine | 7260 | I |