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**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.33(a), this is notice that on October 7, 2022, Sterling Pharma USA, LLC, 1001 Sheldon Drive, Suite 101, Cary, North Carolina 27513-2078, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
5-Methoxy-N-N-dimethyltryptamine.	7431	I
Psilocybin .....	7437	I
Psilocyn .....	7438	I

The company plans to manufacture the above-listed controlled substances as clinical trials. No other activities for these drug codes are authorized for this registration.

**Matthew Strait,**  
Deputy Assistant Administrator.  
[FR Doc. 2022-26913 Filed 12-9-22; 8:45 am]  
**BILLING CODE P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**  
[Docket No. DEA-1119]

**Bulk Manufacturer of Controlled Substances Application: Scottsdale Research Institute**

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

**ACTION:** Notice of application.

**SUMMARY:** Scottsdale Research 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 February 10, 2023. Such persons may also file a written request for a hearing on the application on or before February 10, 2023.

**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 September 28, 2022, Scottsdale Research Institute, 12815 North Cave Creek Road, Phoenix, Arizona 85022, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Psilocybin .....	7437	I
Psilocyn .....	7438	I

The company plans to bulk manufacture the listed controlled substances for internal research purposes and to support clinical trials. No other activities for these drug codes are authorized for this registration.

**Matthew Strait,**  
Deputy Assistant Administrator.  
[FR Doc. 2022-26922 Filed 12-9-22; 8:45 am]  
**BILLING CODE P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**  
[Docket No. DEA-1114]

**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 **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 January 11, 2023. Such persons may also file a written request for a hearing on the application on or before January 11, 2023.

**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 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 September 12, 2022, VHG Labs dba LGC Standards, 3 Perimeter Road, Manchester, New Hampshire 03103-3341, 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