## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

**[Docket No. DEA–1685]**

### Importer of Controlled Substances Application: Patheon API Services, Inc.

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

**ACTION:** Notice of application.

**SUMMARY:** Patheon API Services, Inc. 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 April 20, 2026. Such persons may also file a written request for a hearing on the application on or before April 20, 2026.

**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 19, 2026, Patheon API Services, Inc., 101 Technology Place, Florence, South Carolina 29501, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

| Controlled substance | Drug code | Schedule |
|---|---|---|
| Dimethyltryptamine | 7435 | I |
| Psilocybin | 7437 | I |
| Psilocyn | 7438 | I |
| Amphetamine | 1100 | II |
| Methadone | 9250 | II |

The company plans to import the listed controlled substances as reference standards for research and development as part of API Manufacturing. 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.

**Thomas Prevoznik,**

*Deputy Assistant Administrator.*

[FR Doc. 2026–05355 Filed 3–18–26; 8:45 am]

**BILLING P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

**[Docket No. DEA–1682]**

### Bulk Manufacturer of Controlled Substances Application: Siegfried Grafton, Inc.

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

**ACTION:** Notice of application.

**SUMMARY:** Siegfried Grafton, Inc. 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 May 18, 2026. Such persons may also file a written request for a hearing on the application on or before May 18, 2026.

**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 February 25, 2026, Siegfried Grafton, Inc., 870 Badger Circle, Grafton, Wisconsin 53024–0000, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

| Controlled substance | Drug code | Schedule |
|---|---|---|
| Lysergic acid diethylamide. | 7315 | I |
| Marihuana Extract | 7350 | I |
| Marihuana | 7360 | I |
| Tetrahydrocannabinols | 7370 | I |
| 4-Bromo-2,5-dimethoxy-phenethylamine. | 7392 | I |
| 3,4-Methylenedioxy-amphetamine. | 7400 | I |
| 3,4-Methylenedioxy-methamphetamine. | 7405 | I |
| 5-Methoxy-N-N-dimethyltryptamine. | 7431 | I |
| Dimethyltryptamine | 7435 | I |
| Psilocybin | 7437 | I |
| Psilocyn | 7438 | I |
| Lisdexamfetamine | 1205 | II |
| Methylphenidate | 1724 | II |
| Amobarbital | 2125 | II |
| Nabilone | 7379 | II |
| ANPP (4-Anilino-N-phenethyl-4-piperidine). | 8333 | II |
| Hydrocodone | 9193 | II |
| Opium extracts | 9610 | II |
| Opium, powdered | 9639 | II |
| Opium, granulated | 9640 | II |
| Opium poppy | 9650 | II |
| Noroxymorphone | 9668 | II |
| Remifentanil | 9739 | II |
| Fentanyl | 9801 | II |

The company plans to bulk manufacture the listed controlled substances for purpose of analytical reference standards or for sale to its customers. In reference to drug codes 7360 (Marihuana), and 7370 (Tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetic. No other activities for these drug codes are authorized for this registration.

**Thomas Prevoznik,**

*Deputy Assistant Administrator.*

[FR Doc. 2026–05358 Filed 3–18–26; 8:45 am]

**BILLING CODE 4410–09–P**