

**DEPARTMENT OF JUSTICE****Drug Enforcement Administration****[Docket No. DEA-1653]****Importer of Controlled Substances  
Application: Mylan Inc.****AGENCY:** Drug Enforcement  
Administration, Justice.**ACTION:** Notice of application.

**SUMMARY:** Mylan 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 March 4, 2026. Such persons may also file a written request for a hearing on the application on or before March 4, 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 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 December 19, 2025, Mylan Inc., 3711 Collins Ferry Road, Morgantown, West Virginia 26505-2362, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

| Controlled substance  | Drug code | Schedule |
|-----------------------|-----------|----------|
| Amphetamine .....     | 1100      | II       |
| Methylphenidate ..... | 1724      | II       |
| Oxycodone .....       | 9143      | II       |
| Hydromorphone .....   | 9150      | II       |
| Methadone .....       | 9250      | II       |
| Morphine .....        | 9300      | II       |
| Fentanyl .....        | 9801      | II       |

The company plans to import the listed controlled substances as bulk active pharmaceutical ingredients for internal testing purposes only and finished dosage forms for analytical testing and distribution for clinical trials to support foreign market participation. 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. 2026-01947 Filed 1-30-26; 8:45 am]

**BILLING CODE P****DEPARTMENT OF JUSTICE****Drug Enforcement Administration****[Docket No. DEA-1649]****Bulk Manufacturer of Controlled  
Substances Application: Promega  
Corporation****AGENCY:** Drug Enforcement  
Administration, Justice.**ACTION:** Notice of application.

**SUMMARY:** Promega Corporation 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 April 3, 2026. Such persons may also file a written request for a hearing on the application on or before April 3, 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 January 7, 2026, Promega Corporation, 3075 Sub Zero Parkway, Fitchburg, Wisconsin 53719, 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 as Active Pharmaceutical Ingredients for sale to its customers. No other activities for these drug codes are authorized for this registration.

**Thomas Prevoznik,***Deputy Assistant Administrator.*

[FR Doc. 2026-01940 Filed 1-30-26; 8:45 am]

**BILLING CODE P****DEPARTMENT OF JUSTICE****Drug Enforcement Administration****[Docket No. DEA-1651]****Importer of Controlled Substances  
Application: Sharp Clinical Services,  
LLC****AGENCY:** Drug Enforcement  
Administration, Justice.**ACTION:** Notice of application.

**SUMMARY:** Sharp Clinical Services, LLC 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 March 4, 2026. Such persons may also file a written request for a hearing on the application on or before March 4, 2026.