

**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 January 9, 2026, Sharp Clinical Services, LLC, 2400 Baglyos Circle, Bethlehem, Pennsylvania 18020-8024, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid.	2010	I
3,4-Methylenedioxy-methamphetamine.	7405	I
5-Methoxy-N-N-dimethyltryptamine.	7431	I
Psilocybin .....	7437	I

The company plans to import the listed controlled substances for distribution and clinical trials. 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. 2026-01944 Filed 1-30-26; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-1652]

**Importer of Controlled Substances Application: Medi-Physics Inc. DBA GE Healthcare**

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

**ACTION:** Notice of application.

**SUMMARY:** Medi-Physics Inc. DBA GE Healthcare 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.

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8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on December 17, 2025, Medi-Physics Inc. DBA GE Healthcare, 3350 North Ridge Avenue, Arlington Heights, Illinois 60004-1412, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Cocaine .....	9041	II

The company plans to import derivatives of the listed controlled substance to be used for the manufacture of a diagnostic product and reference standards. No other activity for this drug code 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.

**Thomas Prevoznik,**  
Deputy Assistant Administrator.  
[FR Doc. 2026-01946 Filed 1-30-26; 8:45 am]

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**OFFICE OF THE FEDERAL REGISTER**

**Publication Procedures for Federal Register Documents During a Funding Hiatus**

**AGENCY:** Office of the Federal Register.  
**ACTION:** Notice of special procedures.

**SUMMARY:** During an appropriations lapse, the Office of the Federal Register (OFR) publishes documents that meet an exception under the Antideficiency Act (ADA). It is the responsibility of the agency submitting a document for publication during an appropriations lapse to provide an exception letter with the document that includes a justification and a certification that the document is authorized under an exception to the Antideficiency Act.

**FOR FURTHER INFORMATION CONTACT:** Liza Davis, Esq., Director of Legal Affairs and Policy, Office of the Federal Register, National Archives and Records Administration, (202) 741-6030 or [Fedreg.legal@nara.gov](mailto:Fedreg.legal@nara.gov).

**SUPPLEMENTARY INFORMATION:** In accordance with the provisions of the