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 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-meth-amphetamine.	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]

BILLING CODE P

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.

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 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]

BILLING CODE P

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.*

SUPPLEMENTARY INFORMATION: In accordance with the provisions of the

Antideficiency Act (ADA), Public Law 97–258, as amended (31 U.S.C. 1341, 1342), and Office of Legal Counsel (OLC) Opinions *Government Operations in the Event of a Lapse in Appropriations* (19 Op. O.L.C. 301, August 16, 1995), and *Effect of Appropriations for Other Agencies and Branches on the Authority to Continue Department of Justice Functions During the Lapse in the Department's Appropriations* (19 Op. O.L.C. 337, December 13, 1995), the OFR announces special procedures for agencies transmitting documents for publication in the **Federal Register** during a lapse in appropriations.

During an appropriations lapse, the OFR remains open to accept and process documents authorized to be published in the **Federal Register** in the absence of continuing appropriations. An agency wishing to transmit a document to the OFR during an appropriations lapse must attach an exception letter to the document which provides a justification and certifies that publication in the **Federal Register** is necessary. The OFR will only publish documents submitted during an appropriations lapse that meet an exception to the ADA, with sufficient justification that the document meets the ADA exception as provided by the publishing agency. This may include documents that directly relate to the performance of governmental functions necessary to address imminent threats to the safety of human life or protection of property (the ADA emergency exception) or that meet another exception to the ADA, as well as documents related to funded programs if delaying publication until the end of the appropriations lapse would prevent or significantly damage the execution of funded functions at the agency. It is the responsibility of the agency submitting a document for publication to include an exception letter that provides justification and certifies that the document is authorized under the ADA; the OFR does not provide this justification for the submitting agency. This certification provides OFR with documentation that publication in the **Federal Register** is a function or service excepted under the ADA.

Executive branch agencies and offices should use the template for the exception letter available on the OFR website at *www.archives.gov/federal-register/agencies/shutdown-faqs.* Legislative and judicial branch offices may use the template letter as a guide.

Special handling requests should be included in the exception letter. Do not submit two separate letters.

Documents received and scheduled for publication before the appropriations lapse began are not required to meet an ADA exception.

For final rule documents that contain incorporation by reference (IBR), agencies must submit a separate request for IBR approval as per normal procedure, and must include sufficient justification that the rule document meets an exception to the ADA when submitting the IBR request. The OFR will not review an IBR request that does not include a sufficient justification. Requests without a sufficient justification will be held until the appropriations lapse is ended.

The OFR may suspend the regular publication schedule during an appropriations lapse to permit a limited number of excepted personnel to process excepted documents. Agency officials will be informed as to the schedule for filing and publishing individual documents.

The OFR has posted frequently asked questions and the excepted letter template on the following website: *www.archives.gov/federal-register/agencies/shutdown-faqs.*

Authority: 44 U.S.C. 1502; 1 CFR 2.4 and 5.1.

Liza Davis,

Director of Legal Affairs and Policy, Office of the Federal Register.

[FR Doc. 2026–02066 Filed 1–30–26; 8:45 am]

BILLING CODE 0099–10–P

NATIONAL ARCHIVES AND RECORDS ADMINISTRATION

[NARA–26–0034; NARA–2026–006]

Records Schedules; Availability and Request for Comments

AGENCY: National Archives and Records Administration (NARA).

ACTION: Notice of availability of proposed records schedules; request for comments.

SUMMARY: The National Archives and Records Administration (NARA) publishes notice of certain Federal agency requests for records disposition authority (records schedules). We publish notice in the **Federal Register** and on *regulations.gov* for records schedules in which agencies propose to dispose of records they no longer need to conduct agency business. We invite public comments on such records schedules.

DATES: We must receive responses on the schedules listed in this notice by March 19, 2026.

ADDRESSES: To view a records schedule in this notice, or submit a comment on one, use the following address: *https://www.regulations.gov/docket/NARA-26-0034/document.*

This is a direct link to the schedules posted in the docket for this notice on *regulations.gov.* You may submit comments by the following method:

- *Federal eRulemaking Portal: https://www.regulations.gov.* On the website, enter either of the numbers cited at the top of this notice into the search field. This will bring you to the docket for this notice, in which we have posted the records schedules open for comment. Each schedule has a 'comment' button so you can comment on that specific schedule. For more information on *regulations.gov* and on submitting comments, see their FAQs at *https://www.regulations.gov/faq.*

If you are unable to comment via *regulations.gov,* you may email us at *request.schedule@nara.gov* for instructions on submitting your comment. You must cite the control number of the schedule you wish to comment on. You can find the control number for each schedule in parentheses at the end of each schedule's entry in the list at the end of this notice.

FOR FURTHER INFORMATION CONTACT: Richard Green, Records Management Operations, by email at *richard.green@nara.gov* or at 301–395–7825. For information about records schedules, contact Records Management Operations by email at *request.schedule@nara.gov* or by phone at 301–395–7825.

SUPPLEMENTARY INFORMATION:

Public Comment Procedures

We are publishing notice of records schedules in which agencies propose to dispose of records they no longer need to conduct agency business. We invite public comments on these records schedules, as required by 44 U.S.C. 3303a(a), and list the schedules at the end of this notice by agency and subdivision requesting disposition authority.

In addition, this notice lists the organizational unit(s) accumulating the records or states that the schedule has agency-wide applicability. It also provides the control number assigned to each schedule, which you will need if you submit comments on that schedule. We have uploaded the records schedules and accompanying appraisal memoranda to the *regulations.gov* docket for this notice as ''other'' documents. Each records schedule contains a full description of the records