information from or copies of documents in military personnel, military medical, and dependent medical records. We invite you to comment on the proposed information collections. There have been no changes made to these collections.

DATES: OMB must receive written comments on or before September 10, 2026.

ADDRESSES: Send any comments and recommendations on the proposed information collection in writing to *www.reginfo.gov/public/do/PRAMain.* You can find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

FOR FURTHER INFORMATION CONTACT: Kristin Phillips, Paperwork Reduction Act Officer, by email at *kristin.phillips@nara.gov* or by telephone at 616–254–0405 with any requests for additional information.

SUPPLEMENTARY INFORMATION: Pursuant to the Paperwork Reduction Act of 1995 (Pub. L. 104–13), we invite the public and other Federal agencies to comment on proposed information collections. We published a notice of proposed collection for these information collections on June 1, 2026 (91 FR 32455) and we received no comments. We are therefore submitting the described information collections to OMB for approval.

If you have comments or suggestions, they should address one or more of the following points: (a) whether the proposed information collections are necessary for NARA to properly perform its functions; (b) our estimate of the burden of the proposed information collections and its accuracy; (c) ways we could enhance the quality, utility, and clarity of the information we collect; (d) ways we could minimize the burden on respondents of collecting the information, including through information technology; and (e) whether these collections affect small businesses.

In this notice, we solicit comments concerning the following information collections:

1. *Title:* Request to Digitize Records.
OMB number: 3095–0017.
Agency form number: None.
Type of review: Regular.
Affected public: Companies and organizations that wish to digitize archival holdings in the National Archives of the United States or a Presidential library for micropublication.
Estimated number of respondents: 10.
Estimated time per response: 5 hours.
Frequency of response: On occasion (when respondent wishes to request permission to digitize records).
Estimated total annual burden hours: 50.
Abstract: The information collection is prescribed by 36 CFR 1254.92. The collection is prepared by companies and organizations that wish to digitize archival holdings with privately-owned equipment. NARA uses the information to determine whether the request meets the criteria in 36 CFR 1254.100, to evaluate the records for digitization, and to schedule use of the limited space available for digitizing.

2. *Title:* Forms Relating to Military Service Records.
OMB number: 3095–0039.
Agency form number: NA Forms 13036, 13042, 13055, 13075, and 13177.
Type of review: Regular.
Affected public: Veterans, their authorized representatives, state and local governments, and businesses.
Estimated number of respondents: 79,800.
Estimated time per response: 5 minutes.
Frequency of response: On occasion (when respondent wishes to request information from a military personnel, military medical, and dependent medical record).
Estimated total annual burden hours: 6,650 hours.
Abstract: In accordance with rules issued by the Department of Defense (DOD) and the Department of Homeland Security (DHS, U.S. Coast Guard), the National Personnel Records Center (NPRC) of the National Archives and Records Administration (NARA) administers military personnel and medical records of veterans after discharge, retirement, and death. In addition, NPRC administers the medical records of dependents of service personnel. When veterans, dependents, and other authorized individuals request information from or copies of documents in military personnel, military medical, and dependent medical records, they must provide on forms or in letters certain information about the veteran and the nature of the request. A major fire at the NPRC on July 12, 1973, destroyed numerous military records. If individuals' requests involve records or information from records that may have been lost in the fire, requesters may be asked to complete NA Form 13075, Questionnaire about Military Service, or NA Form 13055, Request for Information Needed to Reconstruct Medical Data, so that NPRC staff can search alternative sources to reconstruct the requested information. Requesters who ask for medical records of dependents of service personnel and hospitalization records of military personnel are asked to complete NA Form 13042, Request for Information Needed to Locate Medical Records, so that NPRC staff can locate the desired records. Certain types of information contained in military personnel and medical records are restricted from disclosure unless the veteran provides a more specific release authorization than is normally required. Veterans are asked to complete NA Form 13036, Authorization for Release of Military Medical Patient Records, to authorize release to a third party of a restricted type of information found in the desired record. For those who have already made a request, and want to check the status, they can use NA Form 13177, Check the Status of a Clinical & Medical Treatment Records Request.

Gulam Shakir,
Executive for Information Services/CIO.
[FR Doc. 2026–16352 Filed 8–10–26; 8:45 am]
BILLING CODE 7515–01–P

NUCLEAR REGULATORY COMMISSION

[NRC–2026–3730]

Draft Regulatory Guide: Release of Patients Administered Radioactive Material

AGENCY: Nuclear Regulatory Commission.

ACTION: Draft guide; request for comment.

SUMMARY: The U.S. Nuclear Regulatory Commission (NRC) is issuing for public comment a draft Regulatory Guide (DG), DG–8070, ''Release of Patients Administered Radioactive Material.'' This DG is proposed Revision 2 of Regulatory Guide (RG) 8.39 of the same name. The proposed revision provides methods that are acceptable to the NRC staff for the release of individuals, referred to as patients in this revision, after a medical procedure involving the administration of unsealed byproduct material, such as radiopharmaceuticals, or implants that contain radioactive material.

DATES: Comments must be submitted electronically using *https://www.regulations.gov* by 11:59 p.m. Eastern Time on September 10, 2026.

ADDRESSES: Submit your comments, identified by Docket ID NRC–2026–3730, at *https://www.regulations.gov.* If your material cannot be submitted using *https://www.regulations.gov,* call or

email the individual listed in the **FOR FURTHER INFORMATION CONTACT** section of this document for alternate instructions.

- Do not include any personally identifiable information (such as name, address, or other contact information) or confidential business information that you do not want publicly disclosed. All comments are public records; they are publicly displayed exactly as received, and will not be deleted, modified, or redacted. Comments may be submitted anonymously.
- Follow the search instructions on *https://www.regulations.gov* to view public comments.
- For additional direction on obtaining information and submitting comments, see ''Obtaining Information and Submitting Comments'' in the **SUPPLEMENTARY INFORMATION** section of this document.

FOR FURTHER INFORMATION CONTACT: Sarah Spence, telephone: 301–415–3697; email: *Sarah.Spence@nrc.gov,* and Amir Mobasheran, telephone: 301–415–8112; email: *Amir.Mobasheran@nrc.gov.* Both are staff of the Office of Nuclear Material Safety and Safeguards, at the U.S. Nuclear Regulatory Commission, Washington, DC 20555–0001.

SUPPLEMENTARY INFORMATION:

I. Obtaining Information and Submitting Comments

A. Obtaining Information

Please refer to Docket ID NRC–2026–3730 when contacting the NRC about the availability of information for this action. You may obtain publicly available information related to this action by any of the following methods:

- *Federal Rulemaking Website:* Go to *https://www.regulations.gov* and search for Docket ID NRC–2026–3730.
- *NRC's Agencywide Documents Access and Management System (ADAMS):* You may obtain publicly available documents online in the ADAMS Public Documents collection at *https://www.nrc.gov/reading-rm/adams.html.* To begin the search, select ''Begin ADAMS Public Search.'' For problems with ADAMS, please contact the NRC's Public Document Room (PDR) reference staff at 1–800–397–4209, at 301–415–4737, or by email to *PDR.Resource@nrc.gov.* The Release of Patients Administered Radioactive Material is available in ADAMS under Accession No. ML26092A365.
- *NRC's PDR:* The PDR, where you may examine and order copies of publicly available documents, is open by appointment. To make an appointment to visit the PDR, please send an email to *PDR.Resource@nrc.gov* or call 1–800–397–4209 or 301–415–4737, between 8 a.m. and 4 p.m. Eastern Time (ET), Monday through Friday, except Federal holidays.

B. Submitting Comments

Comments must be submitted electronically using *https://www.regulations.gov* no later than 11:59 p.m. ET on September 10, 2026. Please include Docket ID NRC–2026–3730 in your comment submission.

The NRC cautions you not to include identifying or contact information that you do not want to be publicly disclosed in your comment submission. The NRC will post all comment submissions at *https://www.regulations.gov* as well as enter the comment submissions into ADAMS. The NRC does not routinely edit comment submissions to remove identifying or contact information.

If you are requesting or aggregating comments from other persons for submission to the NRC, then you should inform those persons not to include identifying or contact information that they do not want to be publicly disclosed in their comment submission. Your request should state that the NRC does not routinely edit comment submissions to remove such information before making the comment submissions available to the public or entering the comment into ADAMS.

II. Additional Information

The NRC is issuing for public comment a DG in the NRC's ''Regulatory Guide'' series. This series was developed to describe methods that are acceptable to the NRC staff for implementing specific parts of the agency's regulations, to explain techniques that the staff uses in evaluating specific issues or postulated events, and to describe information that the staff needs in its review of applications for permits and licenses.

Proposed RG 8.39 Revision 2, entitled ''Release of Patients Administered Radioactive Material,'' is temporarily identified by its task number, DG–8070.

The existing version of RG 8.39 (Revision 1) was published in April 2020. The NRC previously published for comment a draft update of RG 8.39 (DG–8061), in April 2024, received comments relating to the technical complexity and requests for more examples to be included, and did not thereafter finalize that update. The NRC has now refined a draft revision to RG 8.39 and has included draft revisions in DG–8070 to align with proposed revisions to section 35.75 of title 10 of the *Code of Federal Regulations* (10 CFR) as part of a proposed rule, ''Reforming and Modernizing the NRC's Radiation Protection Framework'' (91 FR 43456; July 15, 2026), that addresses section 5(b) of Executive Order 14300.

The proposed revision provides methods that are acceptable to the NRC staff for the release of individuals, referred to as patients in this revision, after a medical procedure involving the administration of unsealed byproduct material, such as radiopharmaceuticals, or implants that contain radioactive material. More specifically, DG–8070 provides guidance and examples for application of new definitions and dose limits related to medical uses of byproduct material that would be introduced by the referenced proposed rule, including guidance related to doses to caregivers.

Additionally, DG–8070 incorporates up-to-date methodologies based on National Council on Radiation Protection and Measurements (NCRP) Report No. 155, ''Management of Radionuclide Therapy Patients,'' fundamentals for calculating the radiation dose received by members of the public from a patient containing byproduct material. DG–8070 also includes updates to reflect the evolution of the medical use of byproduct material since the last revision of RG 8.39 (Revision 1) was issued. This DG is intended to enable licensees to make more accurate estimates of the timing, circumstances, and risks associated with patient release; public doses from the released patients; and when instructions or records are required.

The staff is not issuing for public comment a separate draft regulatory analysis for this draft regulatory guide. The draft regulatory analysis associated with this draft regulatory guide was issued for public comment with the proposed rule, ''Reforming and Modernizing the NRC's Radiation Protection Framework'' (91 FR 43456). That draft regulatory analysis assesses the value associated with issuing or revising regulatory guides as well as alternative courses of action (ADAMS Accession No. ML26180A025).

III. Backfitting, Forward Fitting, and Issue Finality

The NRC staff may use this RG as a reference in its regulatory processes, such as licensing, inspection, or enforcement. Backfitting, forward fitting, and issue finality considerations do not apply to 10 CFR part 35 licensees and applicants because 10 CFR part 35 does not include backfitting or issue finality provisions, and the forward fitting policy in Management Directive 8.4, ''Management of Backfitting, Forward Fitting, Issue Finality, and

Information Requests,'' does not apply to these licensees.

IV. Submitting Suggestions for Improvement of Regulatory Guides

A member of the public may, at any time, submit suggestions to the NRC for improvement of existing RGs or for the development of new RGs. Suggestions can be submitted on the NRC's public website at *https://www.nrc.gov/reading-rm/doc-collections/reg-guides/contactus.html.* Suggestions will be considered in future updates and enhancements to the ''Regulatory Guide'' series.

V. Executive Order (E.O.) 12866

The Office of Information and Regulatory Affairs determined that this DG is not a significant regulatory action under E.O. 12866.

Authority: 42 U.S.C. 2011 *et seq.*

Dated: August 7, 2026.

For the Nuclear Regulatory Commission.

Nicholee Valentine,

Chief, Guidance and Publication Branch, Division of Guidance, Rulemaking, Economic Analysis, and Technical Editing, Office of Nuclear Material Safety and Safeguards.

[FR Doc. 2026–16344 Filed 8–10–26; 8:45 am]

BILLING CODE 7590–01–P

NUCLEAR REGULATORY COMMISSION

[NRC–2026–0001]

Sunshine Act Meetings

TIME AND DATE: Weeks of August 10, 17, 24, 31, and September 7, 14, 2026. The schedule for Commission meetings is subject to change on short notice. The NRC Commission Meeting Schedule can be found on the internet at: *https://www.nrc.gov/public-involve/public-meetings/schedule.html.*

PLACE: The NRC provides reasonable accommodation to individuals with disabilities where appropriate. If you need a reasonable accommodation to participate in these public meetings or need this meeting notice or the transcript or other information from the public meetings in another format (*e.g.,* braille, large print), please contact the Reasonable Accommodations Resource by email at *Reasonable_Accommodations.Resource@nrc.gov.* Determinations on requests for reasonable accommodation will be made on a case-by-case basis.

STATUS: Public.

Members of the public may request to receive the information in these notices electronically. If you would like to be added to the distribution, please contact the Nuclear Regulatory Commission, Office of the Secretary, Washington, DC 20555, at 301–415–1969, or by email at *Betty.Thweatt@nrc.gov* or *Samantha.Miklaszewski@nrc.gov.*

MATTERS TO BE CONSIDERED:

Week of August 10, 2026

There are no meetings scheduled for the week of August 10, 2026.

Week of August 17, 2026—Tentative

There are no meetings scheduled for the week of August 17, 2026.

Week of August 24, 2026—Tentative

There are no meetings scheduled for the week of August 24, 2026.

Week of August 31, 2026—Tentative

There are no meetings scheduled for the week of August 31, 2026.

Week of September 7, 2026—Tentative

There are no meetings scheduled for the week of September 7, 2026.

Week of September 14, 2026—Tentative

There are no meetings scheduled for the week of September 14, 2026.

CONTACT PERSON FOR MORE INFORMATION: For more information or to verify the status of meetings, contact Wesley Held at 301–287–3591 or via email at *Wesley.Held@nrc.gov.*

The NRC is holding the meetings under the authority of the Government in the Sunshine Act, 5 U.S.C. 552b.

Dated: August 7, 2026.

For the Nuclear Regulatory Commission.

Wesley W. Held,

Policy Coordinator, Office of the Secretary.

[FR Doc. 2026–16364 Filed 8–7–26; 4:15 pm]

BILLING CODE 7590–01–P

NUCLEAR REGULATORY COMMISSION

[NRC–2026–3400]

Application for Amendments to Facility Operating Licenses Involving Proposed No Significant Hazards Consideration Determination and Containing Sensitive Unclassified Non-Safeguards Information and Order Imposing Procedures for Access to Sensitive Unclassified Non-Safeguards Information

AGENCY: Nuclear Regulatory Commission.

ACTION: License amendment request; notice of opportunity to comment, request a hearing, and petition for leave to intervene; order imposing procedures.

SUMMARY: The U.S. Nuclear Regulatory Commission (NRC, the Commission) received, and is considering approval of, one request to amend two licenses. The license amendment request is for St. Lucie Plant, Unit Nos. 1 and 2. For the license amendment request, the NRC proposes to determine that it involves no significant hazards consideration (NSHC). Because the amendment request contains sensitive unclassified non-safeguards information (SUNSI), the NRC is issuing an order imposing procedures to obtain access to SUNSI for contention preparation by persons who file a hearing request or petition for leave to intervene.

DATES: Comments must be filed by September 10, 2026. A request for a hearing or petitions for leave to intervene must be filed by October 13, 2026. Any potential party as defined in section 2.4 of title 10 of the *Code of Federal Regulations* (10 CFR) who believes access to SUNSI is necessary to respond to this notice must request document access by August 21, 2026.

ADDRESSES: You may submit comments by any of the following methods; however, the NRC encourages electronic comment submission through the Federal rulemaking website.

- *Federal rulemaking website:* Go to *https://www.regulations.gov* and search for Docket ID NRC–2026–3400. Address questions about Docket IDs in *Regulations.gov* to Bridget Curran; telephone: 301–415–1003; email: *Bridget.Curran@nrc.gov.* For technical questions, contact the individual listed in the **FOR FURTHER INFORMATION CONTACT** section of this document.
- *Mail comments to:* Office of Nuclear Material Safety and Safeguards, Mail Stop: TWFN–5–A85, U.S. Nuclear Regulatory Commission, Washington, DC 20555–0001, ATTN: Program Management, Announcements and Editing Staff.

For additional direction on obtaining information and submitting comments, see ''Obtaining Information and Submitting Comments'' in the **SUPPLEMENTARY INFORMATION** section of this document.

FOR FURTHER INFORMATION CONTACT: Paula Blechman, Office of Nuclear Reactor Regulation, U.S. Nuclear Regulatory Commission, Washington, DC 20555–0001; telephone: 301–415–2242; email: *Paula.Blechman@nrc.gov.*

SUPPLEMENTARY INFORMATION:

I. Obtaining Information and Submitting Comments

A. Obtaining Information

Please refer to Docket ID NRC–2026–3400, facility name, unit number(s),