205–2781. Copies of non-confidential documents filed in connection with this investigation may be viewed on the Commission's electronic docket (EDIS) at *https://edis.usitc.gov.* For help accessing EDIS, please email *EDIS3Help@usitc.gov.* General information concerning the Commission may also be obtained by accessing its internet server at *https://www.usitc.gov.* Hearing-impaired persons are advised that information on this matter can be obtained by contacting the Commission's TDD terminal on (202) 205–1810.

SUPPLEMENTARY INFORMATION: Section 337 of the Tariff Act of 1930 provides that, if the Commission finds a violation, it shall exclude the articles concerned from the United States unless, after considering the effect of such exclusion upon the public health and welfare, competitive conditions in the United States economy, the production of like or directly competitive articles in the United States, and United States consumers, it finds that such articles should not be excluded from entry. (19 U.S.C. 1337(d)(1)). A similar provision applies to cease and desist orders. (19 U.S.C. 1337(f)(1)).

The Commission is soliciting submissions on public interest issues raised by the recommended relief should the Commission find a violation, specifically: a limited exclusion order for a period of 5–6 years, directed to certain glass substrates for liquid crystal displays and products containing the same that are imported, sold for importation, and/or sold after importation by respondents Caihong Display Devices Co., Ltd. d/b/a Irico Display Devices Co., Ltd. of Xianyang City, China (''Caihong Display''); TCL China Star Optoelectronics, Technology Co., Ltd. of Shenzhen City, China (''CSOT''); Xianyang CaiHong Optoelectronics, Technology Co., Ltd. of Xianyang City, China (''CHOT''); and TTE Technology, Inc., d/b/a TCL North America of Irvine, California (''TCL''); and a cease and desist order directed to TCL. Parties are to file public interest submissions pursuant to 19 CFR 210.50(a)(4).

The Commission is interested in further development of the record on the public interest in this investigation. Accordingly, members of the public and interested government agencies are invited to file submissions of no more than five (5) pages, inclusive of attachments, concerning the public interest in light of the ALJ's Recommended Determination on Remedy and Bonding issued in this investigation on August 6, 2026. Comments should address whether issuance of the recommended remedial orders in this investigation, should the Commission find a violation, would affect the public health and welfare in the United States, competitive conditions in the United States economy, the production of like or directly competitive articles in the United States, or United States consumers.

In particular, the Commission is interested in comments that:

(i) explain how the articles potentially subject to the recommended remedial orders are used in the United States;

(ii) identify any public health, safety, or welfare concerns in the United States relating to the recommended orders;

(iii) identify like or directly competitive articles that complainant, its licensees, or third parties make in the United States which could replace the subject articles if they were to be excluded;

(iv) indicate whether complainant, complainant's licensees, and/or third-party suppliers have the capacity to replace the volume of articles potentially subject to the recommended orders within a commercially reasonable time; and

(v) explain how the recommended orders would impact consumers in the United States.

Written submissions must be filed no later than by close of business on September 8, 2026.

Persons filing written submissions must file the original document electronically on or before the deadlines stated above pursuant to 19 CFR 210.4(f). Submissions should refer to the investigation number (''Inv. No. 337–TA–1433'') in a prominent place on the cover page and/or the first page. (*See* Handbook for Electronic Filing Procedures, *https://www.usitc.gov/secretary/documents/handbook_on_filing_procedures.pdf*). Persons with questions regarding filing should contact the Secretary (202–205–2000).

Any person desiring to submit a document to the Commission in confidence must request confidential treatment by marking each document with a header indicating that the document contains confidential information. This marking will be deemed to satisfy the request procedure set forth in Rules 201.6(b) and 210.5(e)(2) (19 CFR 201.6(b), 210.5(e)(2)). Documents for which confidential treatment by the Commission is properly sought will be treated accordingly. Any non-party wishing to submit comments containing confidential information must serve those comments on the parties to the investigation pursuant to the applicable Administrative Protective Order. A redacted non-confidential version of the document must also be filed simultaneously with any confidential filing and must be served in accordance with Commission Rule 210.4(f)(7)(ii)(A) (19 CFR 210.4(f)(7)(ii)(A)). All information, including confidential business information and documents for which confidential treatment is properly sought, submitted to the Commission for purposes of this investigation may be disclosed to and used: (i) by the Commission, its employees and Offices, and contract personnel (a) for developing or maintaining the records of this or a related proceeding, or (b) in internal investigations, audits, reviews, and evaluations relating to the programs, personnel, and operations of the Commission including under 5 U.S.C. Appendix 3; or (ii) by U.S. government employees and contract personnel, solely for cybersecurity purposes. All contract personnel will sign appropriate nondisclosure agreements. All nonconfidential written submissions will be available for public inspection on EDIS.

This action is taken under the authority of section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and in Part 210 of the Commission's Rules of Practice and Procedure (19 CFR part 210).

By order of the Commission.

Issued: August 7, 2026.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2026–16332 Filed 8–10–26; 8:45 am]

BILLING CODE 7020–02–P

NATIONAL ARCHIVES AND RECORDS ADMINISTRATION

[NARA–2026–034]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: National Archives and Records Administration (NARA).

ACTION: Notice.

SUMMARY: We have submitted a request to the Office of Management and Budget (OMB) for approval to continue to use two information collections. The first information collection is prepared by companies and organizations to request to digitize archival holdings with privately-owned equipment. The second information collection is used when veterans, dependents, and other authorized individuals request

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]
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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