90 FR 1754, as amended by Memorandums 25–02 [5] and 25–03 [6] (the ''Universal Notice''). Section III.A.6.a. of the Universal Notice does not require the use of citizen advisory committees but instead encourages their use as a means to gather feedback and recommendations from the public at least annually.

The August 2019 Notice requires that, following HUD's approval of the Action Plan, each grantee form one or more citizen advisory committees to meet in an open forum at least twice a year. The primary function of these groups is to provide increased transparency to the public on use of the CDBG–MIT funds by soliciting and responding to public comments and serving as an on-going public forum to continuously provide input on the CDBG–MIT funded projects and programs. Citizen advisory committees could also be a part of the process to prepare the initial CDBG–MIT Action Plan submitted to HUD, although not expressly required.

The programs outlined in the County's CDBG–MIT Action Plan are tied to hazard mitigation projects identified in their 2020 Multi-Hazard Mitigation Plan (MHMP) which were informed by public outreach and will continue to receive stakeholder input on planned maintenance activities. Furthermore, given the small grant size, the grantee believes the resources required to convene and manage a citizen advisory committee would be better applied toward project implementation and targeted outreach methods on a project basis. Together, these factors indicate that the grantee's request is reasonable given that the CDBG–MIT grant will be used to pursue the implementation of projects outlined in the County's 2020 MHMP and that the approval of this waiver will reduce administrative burden, save financial resources, and streamline recovery efforts.

After reviewing Hawaii County's request and based on the good cause provided herein, the Department is waiving the alternative requirement found in the second paragraph of Section V.A.3.c. of the August 2019 Notice. HUD notes the County's administrative and financial burden of forming a citizen advisory committee for its CDBG–MIT grant and, in alignment with Section III.A.6.a. of the Universal Notice, waives this requirement. As an alternative requirement the grantee is subject to the requirement in section III.A.6.a. of the Universal Notice that states, ''HUD encourages grantees to gather feedback and recommendations on key program decisions from a citizen advisory group at least annually.'' If the County decides later that they do wish to work with a citizen advisory committee, the County can determine the cadence of the meetings and how the group would provide feedback to the grantee. This waiver applies to the second paragraph of Section V.A.3.c. of the August 2019 Notice as it relates to citizen advisory committees for CDBG–MIT funds under Public Law 116–20 for Hawaii County *only.* The County must continue to follow the first paragraph of Section V.A.3.c. as it relates to the availability and accessibility of the Action Plan, substantial amendments, and performance reports, as well as other citizen participation requirements in Section V.A.3.

V. Finding of No Significant Impact

A Finding of No Significant Impact (FONSI) with respect to the environment has been made in accordance with HUD regulations at 24 CFR part 50, which implement section 102(2)(C) of the National Environmental Policy Act of 1969 (42 U.S.C. 4332(2)(C)). The FONSI is available online through the Federal eRulemaking Portal at *http://www.regulations.gov* and HUD's CDBG–DR website at *https://www.hud.gov/hud-partners/community-cdbg-dr-regulations*.

Ronald J. Kurtz,

Assistant Secretary for Community Planning and Development.

[FR Doc. 2026–23696 Filed 11–19–26; 8:45 am]

BILLING CODE 4210–67–P

[5] Memorandum 25–02, ''Revisions made to the Common Application, Waivers, and Alternative Requirements for Community Development Block Grant Disaster Recovery Grantees: The Universal Notice published in the **Federal Register** (90 FR 1754) and Clarifications to the Allocations for Community Development Block Grant Disaster Recovery and Implementation of the CDBG–DR Consolidated Waivers and Alternative Requirements Notice Published in the **Federal Register** (90 FR 4759),'' published on March 19, 2025. Found at *https://www.hud.gov/sites/dfiles/CPD/documents/CPDUniversalnotice.pdf.*

[6] Memorandum 25–03, ''Revisions made to the Common Application, Waivers, and Alternative Requirements for Community Development Block Grant Disaster Recovery Grantees: The Universal Notice published in the **Federal Register** (90 FR 1754),'' published on March 31, 2025. Found at *https://www.hud.gov/sites/default/files/CPD/documents/CDBG-DR/3-31-2025UniversalNoticeChangesMemo.pdf.*

INTERNATIONAL TRADE COMMISSION

Notice of Receipt of Complaint; Solicitation of Comments Relating to the Public Interest

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission has received a complaint entitled *Certain Mammography Systems with Focal Spot Compensation Technology and Components Thereof, Including X-Ray Tubes, DN 3951;* the Commission is soliciting comments on any public interest issues raised by the complaint or complainant's filing pursuant to the Commission's Rules of Practice and Procedure.

FOR FURTHER INFORMATION CONTACT: Lisa R. Barton, Secretary to the Commission, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436, telephone (202) 205–2000. The public version of the complaint can be accessed on the Commission's Electronic Document Information System (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 United States International Trade Commission (USITC) at *https://www.usitc.gov*. The public record for this investigation may be viewed on the Commission's Electronic Document Information System (EDIS) at *https://edis.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: The Commission has received a complaint and a submission pursuant to § 210.8(b) of the Commission's Rules of Practice and Procedure filed on behalf of Hologic, Inc. on November 16, 2026. The complaint alleges violations of section 337 of the Tariff Act of 1930 (19 U.S.C. 1337) in the importation into the United States, the sale for importation, and the sale within the United States after importation of certain mammography systems with focal spot compensation technology and components thereof, including x-ray tubes. The complaint names as a respondent: Siemens Healthineers AG of Germany; Siemens Healthcare GmbH of Germany; and Siemens Medical Solutions USA, Inc. of Malvern, PA. The complainant requests that the Commission issue a limited exclusion order, cease and desist orders and impose a bond upon respondents' alleged infringing articles during the 60-day Presidential review period pursuant to 19 U.S.C. 1337(j).

Proposed respondents, other interested parties, members of the public, and interested government

agencies are invited to file comments on any public interest issues raised by the complaint or § 210.8(b) filing. Comments should address whether issuance of the relief specifically requested by the complainant in this investigation 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 requested remedial orders are used in the United States;

(ii) identify any public health, safety, or welfare concerns in the United States relating to the requested remedial 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 requested exclusion order and/or a cease and desist order within a commercially reasonable time; and

(v) explain how the requested remedial orders would impact United States consumers.

Written submissions on the public interest must be filed no later than by close of business, eight calendar days after the date of publication of this notice in the **Federal Register**. There will be further opportunities for comment on the public interest after the issuance of any final initial determination in this investigation. Any written submissions on other issues must also be filed by no later than the close of business (5:15 p.m.) [1], eight calendar days after publication of this notice in the **Federal Register**. Complainant(s) may file replies to any written submissions no later than three calendar days after the date on which the written submissions were due. Notwithstanding § 201.14(a) of the Commission's Rules of Practice and Procedure, the deadlines for written submissions and replies shall be counted by calendar days starting with the first calendar day following the **Federal Register** publication date or the due date for written submissions, respectively. The provisions of § 201.14(a) regarding the last day of the period still apply concerning the due date of any filings. No other submissions will be accepted, unless requested by the Commission. Any submissions and replies filed in response to this Notice are limited to five (5) pages in length, inclusive of attachments.

Persons filing written submissions must file the original document electronically on or before the deadlines stated above. Submissions should refer to the docket number (''Docket No. 3951'') in a prominent place on the cover page and/or the first page. (*See* Handbook for Electronic Filing Procedures, Electronic Filing Procedures).[2] Please note the Secretary's Office will accept only electronic filings unless an exemption is granted. Filings must be made through the Commission's Electronic Document Information System (EDIS, *https://edis.usitc.gov.*) Persons with questions regarding filing should contact the Secretary at *EDIS3Help@usitc.gov.*

Any person desiring to submit a document to the Commission in confidence must request confidential treatment. All such requests should be directed to the Secretary to the Commission and must include a full statement of the reasons why the Commission should grant such treatment. *See* 19 CFR 201.6. Documents for which confidential treatment by the Commission is properly sought will be treated accordingly. 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,[3] solely for cybersecurity purposes. All nonconfidential written submissions will be available for public inspection at the Office of the Secretary and on EDIS.[4]

This action is taken under the authority of section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and of §§ 201.10 and 210.8(c) of the Commission's Rules of Practice and Procedure (19 CFR 201.10, 210.8(c)).

By order of the Commission.

Issued: November 17, 2026.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2026–23619 Filed 11–19–26; 8:45 am]

BILLING CODE 7020–02–P

[1] *See* 19 CFR 201.3(c).

[2] Handbook for Electronic Filing Procedures: *https://www.usitc.gov/secretary/documents/handbook_on_filing_procedures.pdf.*

[3] All contract personnel will sign appropriate nondisclosure agreements.

[4] Electronic Document Information System (EDIS): *https://edis.usitc.gov.*

INTERNATIONAL TRADE COMMISSION

Notice of Receipt of Complaint; Solicitation of Comments Relating to the Public Interest

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission has received a complaint entitled *Certain Mammography Systems and Components Thereof, DN 3952;* the Commission is soliciting comments on any public interest issues raised by the complaint or complainant's filing pursuant to the Commission's Rules of Practice and Procedure.

FOR FURTHER INFORMATION CONTACT: Lisa R. Barton, Secretary to the Commission, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436, telephone (202) 205–2000. The public version of the complaint can be accessed on the Commission's Electronic Document Information System (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 United States International Trade Commission (USITC) at *https://www.usitc.gov* . The public record for this investigation may be viewed on the Commission's Electronic Document Information System (EDIS) at *https://edis.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: The Commission has received a complaint and a submission pursuant to § 210.8(b) of the Commission's Rules of Practice and Procedure filed on behalf of Siemens Medical Solutions USA, Inc. on November 16, 2026. The complaint alleges violations of section 337 of the Tariff Act of 1930 (19 U.S.C. 1337) in the importation into the United States, the sale for importation, and the sale within the United States after importation of certain mammography systems and components thereof. The complaint names as a respondent: Hologic, Inc. of Marlborough, MA. The