

UNITED STATES INTERNATIONAL TRADE COMMISSION
Washington, D.C.

In the Matter of

**CERTAIN RADIOTHERAPY SYSTEMS
AND TREATMENT PLANNING
SOFTWARE, AND COMPONENTS
THEREOF**

Investigation No. 337-TA-968

**NOTICE OF COMMISSION DETERMINATION TO EXTEND THE DATE FOR
DETERMINING WHETHER TO REVIEW THE FINAL INITIAL DETERMINATION
IN THIS INVESTIGATION AND TO EXTEND THE TARGET DATE FOR
COMPLETION OF THE INVESTIGATION**

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission has determined to extend the date for determining whether to review the final initial determination (“ID”) in this investigation until January 13, 2017, and to extend the target date for completion of this investigation until March 16, 2017.

FOR FURTHER INFORMATION: Ron Traud, Office of the General Counsel, U.S. International Trade Commission, 500 E Street SW., Washington, DC 20436, telephone (202) 205-3427. Copies of non-confidential documents filed in connection with this investigation are or will be available for inspection during official business hours (8:45 a.m. to 5:15 p.m.) in the Office of the Secretary, U.S. International Trade Commission, 500 E Street SW., Washington, DC 20436, telephone 202-205-2000. General information concerning the Commission may also be obtained by accessing its Internet server (<https://www.usitc.gov>). The public record for this investigation may be viewed on the Commission’s electronic docket (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 instituted this investigation on October 30, 2015, based on a complaint filed by Varian Medical Systems, Inc. of Palo Alto, California; and Varian Medical Systems International AG of ZG, Switzerland (collectively, “Varian”). 80 FR 66934 (Oct. 30, 2015). The complaint alleges violations of section 337 of the Tariff Act of 1930, as amended, 19 U.S.C. 1337, in the importation into the United States, the sale for importation, or the sale within the United States after importation of certain radiotherapy systems and treatment planning software, and components thereof by reason of infringement of certain claims of U.S. Patent Nos. 7,945,021 (“the ‘021 patent”); 8,116,430 (“the ‘430 patent”); 8,867,703 (“the ‘703 patent”); 7,880,154 (“the ‘154 patent”); 7,906,770 (“the ‘770 patent”); and

8,696,538 (“the ‘538 patent”). *Id.* The notice of investigation named as respondents Elekta AB of Stockholm, Sweden; Elekta Ltd. of Crawley, United Kingdom; Elekta GmbH of Hamburg, Germany; Elekta Inc. of Atlanta, Georgia; IMPAC Medical Systems, Inc. of Sunnyvale, California; Elekta Instrument (Shanghai) Limited of Shanghai, China; and Elekta Beijing Medical Systems Co. Ltd. of Beijing, China (collectively, “Elekta”). The Office of Unfair Import Investigations (“OUII”) also was named as a party to the investigation. *Id.*

Prior to the evidentiary hearing, Varian withdrew its allegations as to certain patent claims and also added additional claims. *See* Notice of Commission Determination Not to Review an Initial Determination Granting a Motion to Amend the Complaint and Notice of Investigation (Apr. 4, 2016). Varian proceeded at the evidentiary hearing on the following patents and claims: claims 1, 4, 9, and 15 of the ‘021 patent; claims 6 and 18 of the ‘430 patent; claim 1 of the ‘703 patent; claims 23 and 26 of the ‘154 patent; claims 61, 67, and 68 of the ‘770 patent; and claims 26 and 41 of the ‘538 patent.

On October 27, 2016, the ALJ issued the final ID, which finds a violation of section 337 by Elekta as to claims 23 and 26 of the ‘154 patent; claims 26 and 41 of the ‘538 patent; and claim 67 of the ‘770 patent. The ALJ found no violation of section 337 in connection with claim 61 of the ‘770 patent; claims 1, 4, 9, and 15 of the ‘021 patent; claims 6 and 18 of the ‘430 patent; and claim 1 of the ‘703 patent.

The Commission is extending the date for determining whether to review the final ID in this investigation from December 27, 2016, to January 13, 2017. The Commission is also extending the target date for completion of the investigation from February 27, 2017, to March 16, 2017.

The authority for the Commission’s determination is contained in 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.



Lisa R. Barton
Secretary to the Commission

Issued: November 21, 2016