UNITED STATES INTERNATIONAL TRADE COMMISSION
Washington, D.C.

In the Matter of	**Investigation No. 337-TA-1112**
CERTAIN RADIO FREQUENCY MICRO-NEEDLE DERMATOLOGICAL TREATMENT DEVICES AND COMPONENTS THEREOF	

NOTICE OF THE COMMISSION'S DETERMINATION NOT TO REVIEW AN INITIAL DETERMINATION TERMINATING THE INVESTIGATION WITH RESPECT TO FOUR RESPONDENTS

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission has determined not to review an initial determination ("ID") (Order No. 23) issued by the presiding administrative law judge ("ALJ") that terminates the investigation with respect to four respondents.

FOR FURTHER INFORMATION CONTACT: Robert Needham, Office of the General Counsel, U.S. International Trade Commission, 500 E Street, SW., Washington, D.C. 20436, telephone (202) 708-5468. 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, D.C. 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 May 15, 2018, based on a complaint filed by Syneron Medical Ltd. of Yokneam Illit, Israel; Candela Corporation of Wayland, Massachusetts; and General Hospital Corporation d/b/a Massachusetts General Hospital of Boston, Massachusetts (together, "Complainants"). 83 FR 22515-16. The complaint, as supplemented, 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, and the sale within the United States after

importation of certain radio frequency micro-needle dermatological treatment devices and components thereof that infringe one or more claims of U.S. Patent Nos. 9,510,899 and 9,095,357. *Id.* at 22515. The Commission's notice of investigation named numerous respondents, including Ilooda Co., Ltd. of Suwon, Republic of Korea ("Ilooda"); Cutera, Inc. of Brisbane, California ("Cutera"); Emvera Technologies, LLC of Cedartown, Georgia ("Emvera"); and Rohrer Aesthetics, LLC of Homewood, Alabama ("Rohrer"). *Id.* at 22516. The Office of Unfair Import Investigations is not participating in this investigation. *Id.*

On March 28, 2019, Complainants, Ilooda, Cutera, Emvera, and Rohrer filed a joint motion to terminate the investigation with respect to Ilooda, Cutera, Emvera, and Rohrer pursuant to a settlement agreement. No party filed a response to the motion.

On April 9, 2019, the ALJ issued the subject ID, granting the motion pursuant to Commission Rule 210.21(b) (19 CFR 210.21(b)) and terminating the investigation with respect to Ilooda, Cutera, Emvera, and Rohrer. The ALJ found that the motion complied with Rule 210.21(b) and that there is no evidence that termination by settlement has any adverse effect on the public interest. No petitions for review of the ID were filed.

The Commission has determined not to review the subject ID.

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: May 7, 2019