

K895279 LASERMATIC COMBOLASER MODEL 5050Nov 8, 1989
76 days to decisionK895279 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k895279/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Aug 24, 1989
Decision date	Nov 8, 1989
Days to decision	76 days
Third-party review	No

APPLICANT

Company	Lasermatic OY
Location	Finland, FI
Contact	FULLER W MORTON
510(k) history	2 submissions · 1 cleared · 1988-1989

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k895279/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 6, 2026