FDA 510(k) Premarket Notification

K895279 LASERMATIC COMBOLASER MODEL 5050

CLEARED

Nov 8, 1989
76 days to decision

K895279 · Product code: **GEX** · General & Plastic Surgery
Source: 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