CLEARED

K990947 MODIFIED VERSAPULSE SELECT SINGLE WAVELENGTH, HO: YAG AND DUAL WAVELENGTH, HO: YAG/ND: YAG SURGICAL LASERS AND DELIVERY

Apr 27, 1999
36 days to decision

K990947 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k990947/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Mar 22, 1999
Decision date	Apr 27, 1999
Days to decision	36 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Lumenis, Inc.**
Location	Mchenry, IL, US
Contact	MICHELLE DEETON
510(k) history	43 submissions · 43 cleared · 1979-2022

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k990947/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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