CLEARED

K011703 LUMENIS VERSAPULSE POWERSUITE HOLMIUM (HO:YAG) AND DUAL WAVELENGTH (HO:YAG/ND:YAG) SURGICAL LASERS AND DELIVERY DEVICES

Aug 29, 2001
89 days to decision

K011703 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k011703/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jun 1, 2001
Decision date	Aug 29, 2001
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Lumenis**
Location	Pleasanton, CA, US
Contact	LISA MCGRATH
Website	http://www.lumenis.com/
510(k) history	5 submissions · 5 cleared · 2001-2010