

**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 decisionK011703 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k011703/> 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 June 21, 2026