FDA 510(k) Premarket Notification

CLEARED

K043173 Q-SWITCHED ND:YAG LASER TREATMENT HEAD FOR THE LUMENIS QUANTUM SERIES

Jan 31, 2005
76 days to decision

K043173 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k043173/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Nov 16, 2004
Decision date	Jan 31, 2005
Days to decision	76 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Lumenis, Ltd.**
Location	Santa Clara, CA, US
Contact	MARTHA MURARI
Website	http://www.lumenis.com/
510(k) history	28 submissions · 27 cleared · 2003-2021

Lumenis, Ltd. is a global leader in energy-based medical device solutions for aesthetic and vision care. The company develops and commercializes innovative laser, intense pulsed light (IPL), and radiofrequency technologies with a manufacturing facility in Santa Clara, California. Lumenis has received FDA 510(k) clearances from total submissions since 2003. The company specializes in General & Plastic Surgery devices, which represent 93% of its regulatory submissions. Its cleared portfolio includes ablative and fractional CO? laser systems, IPL platforms, and surgical lase...