

K022181 NOVUS VARIA OPHTHALMIC LASER AND DELIVERY DEVICES WITH ACCESSORIES

Oct 1, 2002
90 days to decision

K022181 · Product code: **GEX** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k022181/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jul 3, 2002
Decision date	Oct 1, 2002
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Lumenis, Inc.
Location	Mchenry, IL, US
Contact	KAREN L BAKER
510(k) history	43 submissions · 43 cleared · 1979-2022

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k022181/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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