

K980685 MODIFIED VERSAPULSE SELECT HO:YAG AND HO:YAG/ND:YAG LASERS

May 20, 1998
86 days to decision

K980685 · Product code: **GEX** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k980685/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Feb 23, 1998
Decision date	May 20, 1998
Days to decision	86 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k980685/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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