

K951812 MODIFIED COHERENT ULTRAPULSE CO2 SURGICAL LASERSJul 7, 1995
79 days to decisionK951812 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k951812/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Apr 19, 1995
Decision date	Jul 7, 1995
Days to decision	79 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k951812/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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