

**K972347 MODIFIED COHERENT VERSAPULSE AESTHETIC  
SURGICAL LASER**Sep 19, 1997  
87 days to decisionK972347 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k972347/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jun 24, 1997
Decision date	Sep 19, 1997
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k972347/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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