

**K904229 MULTILASE 60**Dec 12, 1990  
90 days to decisionK904229 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k904229/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Sep 13, 1990
Decision date	Dec 12, 1990
Days to decision	90 days
Third-party review	No

**APPLICANT**

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Company	<b>Laser Photonics, Inc.</b>
Location	Orlando, FL, US
Contact	ROBERT ANSELMO
510(k) history	14 submissions · 14 cleared · 1988-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k904229/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 6, 2026