

**K974142 HM-RUBY**Jan 28, 1998  
86 days to decisionK974142 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k974142/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Nov 3, 1997
Decision date	Jan 28, 1998
Days to decision	86 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>American Lasers, Inc.</b>
Location	Alhambra, CA, US
Contact	DAVID K QUON
510(k) history	5 submissions · 5 cleared · 1995-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k974142/> 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 June 29, 2026