

**K924647 DLR-1 DERMA-LASE RUBY LASER/DLR-1/DERMA-LASE SYST**Jun 4, 1993  
263 days to decisionK924647 · Product code: **GEX** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k924647/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Sep 14, 1992
Decision date	Jun 4, 1993
Days to decision	263 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Derma-Lase Co.</b>
Location	Chicago, IL, US
Contact	MICHAEL L BARRETTI
510(k) history	5 submissions · 5 cleared · 1992-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k924647/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated July 6, 2026