

**K990075 LPULSA SYL-9000 OPHTHALMIC YAG LASER**Mar 30, 1999  
78 days to decisionK990075 · Product code: **HQF** · Ophthalmic  
Source: <https://www.510kdatabase.net/k990075/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Jan 11, 1999
Decision date	Mar 30, 1999
Days to decision	78 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Light-Med(Usa), Inc.</b>
Location	San Clemente, CA, US
Contact	THOMAS V KEELEY
510(k) history	2 submissions · 2 cleared · 1999-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k990075/> 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 28, 2026