

**K951117 LIHTAN 532 LASER**Oct 18, 1995  
222 days to decisionK951117 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k951117/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Mar 10, 1995
Decision date	Oct 18, 1995
Days to decision	222 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Buckman Co., Inc.</b>
Location	Mchenry, IL, US
Contact	DAVID W SCHLERF
510(k) history	111 submissions · 104 cleared · 1983-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k951117/> 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 May 21, 2026