

**K881848 MODIFIED MODEL 2100 ND:YAG LASER**May 11, 1988  
9 days to decisionK881848 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k881848/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	May 2, 1988
Decision date	May 11, 1988
Days to decision	9 days
Third-party review	No

**APPLICANT**

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Company	<b>Sharplan Lasers, Inc.</b>
Location	Allendale, NJ, US
Contact	STEPHEN DALTON
510(k) history	78 submissions · 78 cleared · 1986-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881848/>; 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 20, 2026