

**K883563 SHARPLAN 2100 ND:YAG LASER, SERIES 2900 SYN.
TIPS**Sep 27, 1988
39 days to decisionK883563 · Product code: **GEX** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k883563/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Aug 19, 1988
Decision date	Sep 27, 1988
Days to decision	39 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k883563/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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