

K932156 MULTILINEOct 15, 1993
164 days to decisionK932156 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k932156/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	May 4, 1993
Decision date	Oct 15, 1993
Days to decision	164 days
Third-party review	No
Summary / Statement	Statement
Other names	DLY-1

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k932156/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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