FDA 510(k) Premarket Notification

K932156 MULTILINE

CLEARED

Oct 15, 1993
164 days to decision

K932156 · Product code: **GEX** · General & Plastic Surgery
Source: 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