FDA 510(k) Premarket Notification

K881848 MODIFIED MODEL 2100 ND:YAG LASER

CLEARED

May 11, 1988
9 days to decision

K881848 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k881848/

SUBMISSION DETAILS

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

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