FDA 510(k) Premarket Notification

K904229 MULTILASE 60

CLEARED

Dec 12, 1990
90 days to decision

K904229 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k904229/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Sep 13, 1990
Decision date	Dec 12, 1990
Days to decision	90 days
Third-party review	No

APPLICANT

Company	**Laser Photonics, Inc.**
Location	Orlando, FL, US
Contact	ROBERT ANSELMO
510(k) history	14 submissions · 14 cleared · 1988-1994