FDA 510(k) Premarket Notification

K852860 MODEL 920 ARGON/DYE LASER-FOR DERMATOLOGICAL/USE

CLEARED

Oct 7, 1985
91 days to decision

K852860 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k852860/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jul 8, 1985
Decision date	Oct 7, 1985
Days to decision	91 days
Third-party review	No

APPLICANT

Company	**Coherent Medical Group**
Location	Palo Alto, CA, US
Contact	LEN GOLDFINE
510(k) history	27 submissions · 24 cleared · 1985-1997