

**K852860 MODEL 920 ARGON/DYE LASER-FOR
DERMATOLOGICAL/USE**Oct 7, 1985
91 days to decisionK852860 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k852860/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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