

K895786 ENDOGUIDE PRODUCT LINE: EX0120, EX0300 AND EX0400Oct 27, 1989
32 days to decisionK895786 · Product code: **GEX** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k895786/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Sep 25, 1989
Decision date	Oct 27, 1989
Days to decision	32 days
Third-party review	No

APPLICANT

Company	Luxar Corp.
Location	Bothell, WA, US
Contact	D LAAKMANN
510(k) history	17 submissions · 17 cleared · 1988-1996

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k895786/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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