FDA 510(k) Premarket Notification

K895786 ENDOGUIDE PRODUCT LINE: EX0120, EX0300 AND EX0400

CLEARED

Oct 27, 1989
32 days to decision

K895786 · 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