FDA 510(k) Premarket Notification

K895559 MODIFIED ENDOGUIDE

CLEARED

Oct 20, 1989
49 days to decision

K895559 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k895559/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Sep 1, 1989
Decision date	Oct 20, 1989
Days to decision	49 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