FDA 510(k) Premarket Notification

K893642 MODIFIED ENDOGUIDE

CLEARED

Jul 14, 1989
60 days to decision

K893642 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k893642/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	May 15, 1989
Decision date	Jul 14, 1989
Days to decision	60 days
Third-party review	No

APPLICANT

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