FDA 510(k) Premarket Notification

K890174 SLT CLMD CONTACT LASER SYSTEM (ADDIT. INDICATION)

CLEARED

Jan 26, 1989
17 days to decision

K890174 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k890174/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jan 9, 1989
Decision date	Jan 26, 1989
Days to decision	17 days
Third-party review	No

APPLICANT

Company	**Surgical Laser Technologies, Inc.**
Location	Villa Hills, KY, US
Contact	GIFFORD, R.N.
510(k) history	51 submissions · 51 cleared · 1985-2004