

K890174 SLT CLMD CONTACT LASER SYSTEM (ADDIT. INDICATION)Jan 26, 1989
17 days to decisionK890174 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k890174/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 27, 2026