

**K042211 MODIFICATION TO: LASERPRO 810, 940, AND 980
DIODE DIODE LASER SYSTEMS**Nov 8, 2004
84 days to decisionK042211 · Product code: **GEX** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k042211/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Aug 16, 2004
Decision date	Nov 8, 2004
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k042211/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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