FDA 510(k) Premarket Notification

K042211 MODIFICATION TO: LASERPRO 810, 940, AND 980 DIODE DIODE LASER SYSTEMS

CLEARED

Nov 8, 2004
84 days to decision

K042211 · 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