FDA 510(k) Premarket Notification

K040617 RELIANT LASER SYSTEM II

CLEARED

Jun 15, 2004
98 days to decision

K040617 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k040617/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Mar 9, 2004
Decision date	Jun 15, 2004
Days to decision	98 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Reliant Technologies, Inc.**
Location	Foster City, CA, US
Contact	HEATHER TANNER
510(k) history	19 submissions · 19 cleared · 1994-2008