

K040617 RELIANT LASER SYSTEM IIJun 15, 2004
98 days to decisionK040617 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k040617/> 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 June 25, 2026