FDA 510(k) Premarket Notification

K073000 LUMAPROBE

CLEARED

Oct 1, 2008
343 days to decision

K073000 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k073000/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Oct 24, 2007
Decision date	Oct 1, 2008
Days to decision	343 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Clareblend, Inc.**
Location	Sylmar, CA, US
Contact	JILL CREASY
510(k) history	3 submissions · 3 cleared · 1988-2008