

K073000 LUMAPROBEOct 1, 2008
343 days to decisionK073000 · Product code: **GEX** · General & Plastic SurgerySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k073000/> 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 24, 2026