

K082876 BEAMAX/ FORMAX PULSED LIGHT DEVICE FAMILYNov 10, 2008
42 days to decisionK082876 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k082876/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Sep 29, 2008
Decision date	Nov 10, 2008
Days to decision	42 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Sharplight Technologies, Ltd.
Location	Zichron Yaacov, IL
Contact	ILAN SHARON
510(k) history	4 submissions · 4 cleared · 2008-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k082876/> 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 May 26, 2026