

K080121 CYNOSURE SMARTLIPO MULTIWAVLENGTH LASERJan 31, 2008
14 days to decisionK080121 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k080121/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jan 17, 2008
Decision date	Jan 31, 2008
Days to decision	14 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cynosure, Inc.
Location	Bedford, MA, US
Contact	GEORGE CHO
510(k) history	98 submissions · 98 cleared · 1992-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k080121/> 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 18, 2026