

K990725 NUVOLASE 532 LASER SYSTEMMar 22, 1999
17 days to decisionK990725 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k990725/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Mar 5, 1999
Decision date	Mar 22, 1999
Days to decision	17 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	American Laser Corp.
Location	Mchenry, IL, US
Contact	DANIEL HOFFER
510(k) history	5 submissions · 5 cleared · 1982-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k990725/> 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 21, 2026