

K123443 PRECISE SHP DIODE LASERAug 13, 2013
278 days to decisionK123443 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k123443/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Nov 8, 2012
Decision date	Aug 13, 2013
Days to decision	278 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	CAO Group, Inc.
Location	West Jordan, UT, US
Contact	ROBERT K LARSEN
Website	https://www.caogroup.com
510(k) history	31 submissions · 31 cleared · 2001-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k123443/> 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 8, 2026