

**K131059 PIONEER ELITE DIODE LASER**Sep 13, 2013  
150 days to decisionK131059 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k131059/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Apr 16, 2013
Decision date	Sep 13, 2013
Days to decision	150 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k131059/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 8, 2026