

**K020849 APEX 800 LASER SYSTEM, MODEL APEX 800**May 20, 2002  
66 days to decisionK020849 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k020849/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Mar 15, 2002
Decision date	May 20, 2002
Days to decision	66 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Iridex Corp</b>
Location	Mountain View, CA, US
Contact	JOHN D&apos;ANGELO
510(k) history	13 submissions · 13 cleared · 1998-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020849/> 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 15, 2026