

**K992765 CYNOSURE DIODERM**Sep 8, 1999  
22 days to decisionK992765 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k992765/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Aug 17, 1999
Decision date	Sep 8, 1999
Days to decision	22 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k992765/> 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 18, 2026