

**K912502 LAPAROSCOPIC LASER FIBER DIVERTER**Jul 25, 1991  
50 days to decisionK912502 · Product code: **GCJ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k912502/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, General & Plastic Surgery (GCJ)
Date received	Jun 5, 1991
Decision date	Jul 25, 1991
Days to decision	50 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>United States Endoscopy Group, Inc.</b>
Location	Mentor, OH, US
Contact	LISA M SCHOLZ
510(k) history	94 submissions · 92 cleared · 1991-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k912502/> 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