

**K874680 LASEGUIDE 600A, 600B, 400A, AND 400B**Mar 23, 1988  
131 days to decisionK874680 · Product code: **FCS** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k874680/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Catheter, Fiberoptic, Glass, Ureteral (FCS)
Date received	Nov 13, 1987
Decision date	Mar 23, 1988
Days to decision	131 days
Third-party review	No

**APPLICANT**

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Company	<b>Laser Peripherals, LLC</b>
Location	Hingham, MA, US
Contact	MICHAEL N BASEL
510(k) history	12 submissions · 12 cleared · 1988-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k874680/>, 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 May 27, 2026