

**K880591 LASEGUIDE 600A, 600B, 400A, 400B FOR OB-GYN  
USE**Jun 1, 1988  
111 days to decisionK880591 · Product code: **HHR** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k880591/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Surgical, Gynecologic (HHR)
Date received	Feb 11, 1988
Decision date	Jun 1, 1988
Days to decision	111 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/k880591/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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