

**K122905 PEREGRINE 23GA CURVED LASER PROBE**Apr 30, 2013  
221 days to decisionK122905 · Product code: **HQB** · Ophthalmic  
Source: <https://www.510kdatabase.net/k122905/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Photocoagulator And Accessories (HQB)
Date received	Sep 21, 2012
Decision date	Apr 30, 2013
Days to decision	221 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Peregrine Surgical , Ltd.</b>
Location	Doylestown, PA, US
Contact	RYAN O'LEARY
510(k) history	19 submissions · 19 cleared · 1994-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k122905/> 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 4, 2026