

**K111531 RHA2020**Jul 8, 2011  
36 days to decisionK111531 · Product code: **HLI** · Ophthalmic  
Source: <https://www.510kdatabase.net/k111531/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ophthalmoscope, Ac-powered (HLI)
Date received	Jun 2, 2011
Decision date	Jul 8, 2011
Days to decision	36 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Annidis Health Systems Corp.</b>
Location	Ottawa, Ontario, CA
Contact	MICHAEL MCDONNELL
510(k) history	1 submissions · 1 cleared · 2011-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k111531/> 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 19, 2026