

**K091305 CLEARPATH**Oct 23, 2009  
172 days to decisionK091305 · Product code: **FDF** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k091305/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Colonoscope And Accessories, Flexible/rigid (FDF)
Date received	May 4, 2009
Decision date	Oct 23, 2009
Days to decision	172 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Easyglide , Ltd.</b>
Location	Bonita Springs, FL, US
Contact	PAUL DRYDEN
510(k) history	5 submissions · 5 cleared · 2009-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k091305/> 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 14, 2026