

**K053028 TECHDEVICE GUIDEWIRE**Jan 31, 2006  
96 days to decisionK053028 · Product code: **OCY** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k053028/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscopic Guidewire, Gastroenterology-urology (OCY)
Date received	Oct 27, 2005
Decision date	Jan 31, 2006
Days to decision	96 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Techdevice Corporation</b>
Location	Watertown, MA, US
Contact	LEIGH HAYWARD
510(k) history	4 submissions · 4 cleared · 2006-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k053028/> 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