

**K072354 GUIDEWIRE, MODEL G-V220-3527S AND OTHERS**Dec 7, 2007  
107 days to decisionK072354 · Product code: **OCY** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k072354/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscopic Guidewire, Gastroenterology-urology (OCY)
Date received	Aug 22, 2007
Decision date	Dec 7, 2007
Days to decision	107 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Accellent, Inc.</b>
Location	Wilmington, MA, US
Contact	KARL STEINECK
510(k) history	1 submissions · 1 cleared · 2007-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k072354/> 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 26, 2026