

**K011759 RADIUS NEXT GENERATION GUIDEWIRE**Aug 23, 2001  
78 days to decisionK011759 · Product code: **OCY** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k011759/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscopic Guidewire, Gastroenterology-urology (OCY)
Date received	Jun 6, 2001
Decision date	Aug 23, 2001
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Radius Medical Technologies, Inc.</b>
Location	Hopkinton, MA, US
Contact	MAUREEN FINLAYSON
510(k) history	9 submissions · 9 cleared · 1997-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k011759/> 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