

**K021441 RADIUS SNARE**Jun 14, 2002  
39 days to decisionK021441 · Product code: **MMX** · CardiovascularSource: <https://www.510kdatabase.net/k021441/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Percutaneous Retrieval (MMX)
Date received	May 6, 2002
Decision date	Jun 14, 2002
Days to decision	39 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/k021441/> 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