

**K022813 STEERABLE GUIDEWIRE**Feb 14, 2003  
172 days to decisionK022813 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k022813/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Aug 26, 2002
Decision date	Feb 14, 2003
Days to decision	172 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Lake Region Mfg., Inc.</b>
Location	Mchenry, IL, US
Contact	KAREN MORTENSEN
510(k) history	42 submissions · 42 cleared · 1977-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k022813/> 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 May 22, 2026