

**K140536 HI TORQUE CONNECT GUIDEWIRE, HI TORQUE  
CONNECT FLEX GUIDEWIRE, HI TORQUE CONNECT 250T  
GUIDEWIRE**Dec 17, 2014  
288 days to decisionK140536 · Product code: **DQX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k140536/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Mar 4, 2014
Decision date	Dec 17, 2014
Days to decision	288 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Lake Region Medical</b>
Location	Chaska, MN, US
Contact	KENNETH WALSH
510(k) history	16 submissions · 16 cleared · 2008-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k140536/>; 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 14, 2026