

**K091825 HI-TORQUE PROGRESS GUIDE WIRE FAMILY**Sep 25, 2009  
98 days to decisionK091825 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k091825/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jun 19, 2009
Decision date	Sep 25, 2009
Days to decision	98 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Abbott Vascular, Inc.</b>
Location	Redwood, CA, US
Contact	MICHELE WALZ
510(k) history	20 submissions · 17 cleared · 2000-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k091825/> 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