

**K101648 HI-TORQUE WINN GUIDE WIRE FAMILY**Jul 8, 2010  
27 days to decisionK101648 · Product code: **DQX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k101648/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jun 11, 2010
Decision date	Jul 8, 2010
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k101648/> 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