

**K072460 HI-TORQUE BALANCE MIDDLEWEIGHT UNIVERSAL II  
GUIDE WIRE**Apr 11, 2008  
224 days to decisionK072460 · Product code: **DQX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k072460/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Aug 31, 2007
Decision date	Apr 11, 2008
Days to decision	224 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/k072460/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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