

**K081775 THE MEDCOMP .010 VASCULAR GUIDEWIRES**Mar 25, 2009  
275 days to decisionK081775 · Product code: **DQX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k081775/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jun 23, 2008
Decision date	Mar 25, 2009
Days to decision	275 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medcomp</b>
Location	Mchenry, IL, US
Contact	LISA WEIKERT
510(k) history	40 submissions · 34 cleared · 1982-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k081775/> 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 June 7, 2026