

**K050873 CANALIZER HYDROPHILIC GUIDE WIRE**May 26, 2005  
50 days to decisionK050873 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k050873/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Apr 6, 2005
Decision date	May 26, 2005
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medical Device Technologies, Inc.</b>
Location	Gainesville, FL, US
Contact	KRISTINE LIBERACKI
510(k) history	46 submissions · 46 cleared · 1992-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k050873/> 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 20, 2026