

**K843012 WIREGUIDE OR GUIDEWIRE**Nov 29, 1984  
121 days to decisionK843012 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k843012/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jul 31, 1984
Decision date	Nov 29, 1984
Days to decision	121 days
Third-party review	No

**APPLICANT**

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Company	<b>Medi-Tech, Inc.</b>
Location	Mchenry, IL, US
Contact	SAMUEL D WADE
510(k) history	36 submissions · 35 cleared · 1978-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k843012/>; 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 27, 2026