

**K883880 KATZEN THROMBOLYSIS GUIDEWIRE**Apr 6, 1989  
205 days to decisionK883880 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k883880/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Sep 13, 1988
Decision date	Apr 6, 1989
Days to decision	205 days
Third-party review	No

**APPLICANT**

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Company	<b>Medi-Tech, Inc.</b>
Location	Mchenry, IL, US
Contact	ALBERT SEPRINSKI
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/k883880/>, 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