

**K871850 TELEFLEX STEERABLE GUIDEWIRE**Jan 27, 1988  
260 days to decisionK871850 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k871850/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	May 12, 1987
Decision date	Jan 27, 1988
Days to decision	260 days
Third-party review	No

**APPLICANT**

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Company	<b>Teleflexmedical, Inc.</b>
Location	Jeffrey, NH, US
Contact	RICHARD GARDNER
510(k) history	64 submissions · 61 cleared · 1985-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k871850/>, 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 28, 2026