

**K880761 TAPER-22 STEERABLE GUIDEWIRE**Aug 19, 1988  
176 days to decisionK880761 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k880761/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Feb 25, 1988
Decision date	Aug 19, 1988
Days to decision	176 days
Third-party review	No

**APPLICANT**

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Company	<b>Target Therapeutics</b>
Location	Los Angeles, CA, US
Contact	JONI M SNYDER
510(k) history	70 submissions · 70 cleared · 1985-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880761/>, 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 30, 2026