

**K883579 CORONARY GUIDE WIRE**Nov 4, 1988  
74 days to decisionK883579 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k883579/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Aug 22, 1988
Decision date	Nov 4, 1988
Days to decision	74 days
Third-party review	No

**APPLICANT**

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Company	<b>Datascope Corp.</b>
Location	Mchenry, IL, US
Contact	SCHNEIDER, PHD
510(k) history	136 submissions · 135 cleared · 1976-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k883579/>; 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 20, 2026