

**K870664 GUIDE WIRE**Mar 25, 1987  
34 days to decisionK870664 · Product code: **DQX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k870664/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Feb 19, 1987
Decision date	Mar 25, 1987
Days to decision	34 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/k870664/>; 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