

**K913631 .014 HI-TORQUE FLOPPY PATHFINDER GUIDE WIRE**Oct 28, 1991  
74 days to decisionK913631 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k913631/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Aug 15, 1991
Decision date	Oct 28, 1991
Days to decision	74 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Advanced Cardiovascular Systems, Inc.</b>
Location	Santa Clara, CA, US
Contact	JANE E BEGGS
510(k) history	103 submissions · 100 cleared · 1982-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k913631/> 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 July 1, 2026