

**K920891 ROADRUNNER(TM) WIRE GUIDE, MODIFICATION**Dec 21, 1992  
301 days to decisionK920891 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k920891/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Feb 24, 1992
Decision date	Dec 21, 1992
Days to decision	301 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Cook, Inc.</b>
Location	Mchenry, IL, US
Contact	APRIL LAVENDER
510(k) history	190 submissions · 179 cleared · 1976-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k920891/> 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 May 14, 2026