

**K041762 FINISHING WIRE SUPPORTRAK**Aug 6, 2004  
37 days to decisionK041762 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k041762/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jun 30, 2004
Decision date	Aug 6, 2004
Days to decision	37 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Guidant Corporation</b>
Location	St. Paul, MN, US
Contact	JENNIFER TANG
510(k) history	15 submissions · 12 cleared · 2003-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k041762/> 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