

**K031505 RAPIDO CUT-AWAY GUIDING CATHETER**Jun 25, 2003  
42 days to decisionK031505 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k031505/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Percutaneous (DQY)
Date received	May 14, 2003
Decision date	Jun 25, 2003
Days to decision	42 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Guidant Corporation</b>
Location	St. Paul, MN, US
Contact	SHAH M HILALI
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/k031505/> 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