

**K001694 GUIDANT RETRIEVER DEVICE**May 15, 2001  
347 days to decisionK001694 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k001694/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jun 2, 2000
Decision date	May 15, 2001
Days to decision	347 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Guidant Cardiac and Vascular Surgery</b>
Location	Menlo Park, CA, US
Contact	BERNICE JURIS
510(k) history	7 submissions · 7 cleared · 1997-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k001694/> 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 20, 2026