

**K871021 GUIDING CATHETERS 9F JL4**Jun 16, 1987  
92 days to decisionK871021 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k871021/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Mar 16, 1987
Decision date	Jun 16, 1987
Days to decision	92 days
Third-party review	No

**APPLICANT**

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Company	<b>Devices For Vascular Intervention, Inc.</b>
Location	Redwood City, CA, US
Contact	JAMES F PFEIFFER
510(k) history	14 submissions · 14 cleared · 1987-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k871021/>, 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 6, 2026