

**K992142 SCIMED ANGIOGRAPHIC CATHETERS**Jul 21, 1999  
27 days to decisionK992142 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k992142/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jun 24, 1999
Decision date	Jul 21, 1999
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Scimed</b>
Location	Maple Grove, MN, US
Contact	MELANIE RASKA
510(k) history	8 submissions · 8 cleared · 1990-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k992142/> 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 4, 2026