

**K980973 MEDTRONIC ANGIOGRAPHIC CATHETER**Dec 10, 1998  
269 days to decisionK980973 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k980973/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Mar 16, 1998
Decision date	Dec 10, 1998
Days to decision	269 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronics Interventional Vascular</b>
Location	Danvers, MA, US
Contact	MARK CHARTIER
510(k) history	21 submissions · 21 cleared · 1992-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k980973/> 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 14, 2026