

**K950179 MEDTRONIC(R) GC III CORONARY GUIDING  
CATHETER**Jul 25, 1995  
189 days to decisionK950179 · Product code: **DQY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k950179/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jan 17, 1995
Decision date	Jul 25, 1995
Days to decision	189 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Medtronic Interventional Vascular</b>
Location	Danvers, MA, US
Contact	JOSEPH O MAGLIOZZI
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/k950179/>; 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