

**K854018 ANGIOGRAPHY CATHETERS**Jan 16, 1986  
108 days to decisionK854018 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k854018/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Sep 30, 1985
Decision date	Jan 16, 1986
Days to decision	108 days
Third-party review	No

**APPLICANT**

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Company	<b>Micor, Inc.</b>
Location	Pittsburgh, PA, US
Contact	STEPHEN BRUSHEY
510(k) history	8 submissions · 8 cleared · 1986-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k854018/>; 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 25, 2026