

**K083922 MICRUS MICROCATHETER, COURIER FLEX, MODEL MSF 170000-00**May 6, 2009  
126 days to decisionK083922 · Product code: **DQO** · Cardiovascular  
Source: <https://www.510kdatabase.net/k083922/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Dec 31, 2008
Decision date	May 6, 2009
Days to decision	126 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Micrus Endovascular Corporation</b>
Location	Sunnyvale, CA, US
Contact	PATRICK LEE
510(k) history	23 submissions · 23 cleared · 2005-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k083922/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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