

**K032624 MICRUS MICROCATHETER**Sep 10, 2003  
15 days to decisionK032624 · Product code: **DQO** · Cardiovascular  
Source: <https://www.510kdatabase.net/k032624/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Aug 26, 2003
Decision date	Sep 10, 2003
Days to decision	15 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Micrus Corp.</b>
Location	Mountain View, CA, US
Contact	MARGARET WEBBER
510(k) history	6 submissions · 6 cleared · 2001-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k032624/> 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 June 30, 2026