

**K030201 MODIFICATION TO EXPORT ASPIRATION CATHETER**Mar 26, 2003  
64 days to decisionK030201 · Product code: **NFA** · CardiovascularSource: <https://www.510kdatabase.net/k030201/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Temporary Coronary Saphenous Vein Bypass Graft For Embolic Protection (NFA)
Date received	Jan 21, 2003
Decision date	Mar 26, 2003
Days to decision	64 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronic Vascular</b>
Location	Walker, MI, US
Contact	FRED L BOUCHER
510(k) history	475 submissions · 453 cleared · 1977-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030201/> 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 18, 2026