

**K022272 MALLEABLE SINGLE STAGE VENOUS CANNULA,  
MODELS 68112-68140 (15 TOTAL CODES)**Jan 8, 2003  
177 days to decisionK022272 · Product code: DWF · Cardiovascular  
Source: <https://www.510kdatabase.net/k022272/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Jul 15, 2002
Decision date	Jan 8, 2003
Days to decision	177 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronic Cardiac Surgical Products</b>
Location	Grand Rapids, MI, US
Contact	ROGER BRINK
510(k) history	7 submissions · 7 cleared · 2000-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k022272/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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