

**K013013 MEDTRONIC DLP ARTERIAL CANNULA WITH 3D TIP -  
22 FR.**

Dec 4, 2001  
88 days to decision

K013013 · Product code: **DWF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k013013/>

**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	Sep 7, 2001
Decision date	Dec 4, 2001
Days to decision	88 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 W 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/k013013/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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