

**K031518 EOPA CAP ELONGATED ONE-PIECE ARTERIAL CANNULA WITH PRESSURE MONITORING LINE, MODELS 774XX, 775XX, 776XX, 777XX, 788XX,**

May 22, 2003  
7 days to decision

K031518 · Product code: **DWF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k031518/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	May 15, 2003
Decision date	May 22, 2003
Days to decision	7 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronic Vascular</b>
Location	Walker, MI, US
Contact	DAWN STENSTROM
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/k031518/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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