

**K902355 MCR4000 & MCR4000F CARDIOTOMY RESERVOIR**Aug 23, 1990  
86 days to decisionK902355 · Product code: **DTP** · CardiovascularSource: <https://www.510kdatabase.net/k902355/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Defoamer, Cardiopulmonary Bypass (DTP)
Date received	May 29, 1990
Decision date	Aug 23, 1990
Days to decision	86 days
Third-party review	No

**APPLICANT**

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Company	<b>Medtronic Vascular</b>
Location	Walker, MI, US
Contact	LOWE, PHD
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/k902355/>; 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 14, 2026