

**K090869 MIRCSP (MINIMALLY INVASIVE RETROGRADE  
CORONARY SINUS PERFUSION) MANUAL INFLATE CANNULA,  
MODEL 94113TD**Aug 27, 2009  
149 days to decisionK090869 · Product code: **DWF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k090869/>**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	Mar 31, 2009
Decision date	Aug 27, 2009
Days to decision	149 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronic Perfusion Systems</b>
Location	Minneapolis, MN, US
Contact	JESSICA SIXBERRY
510(k) history	29 submissions · 29 cleared · 2000-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090869/>; 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