

**K924642 MEDTRONIC BIO-MEDICUS FEMORAL  
CANNULA/INTRODUCER**Mar 26, 1993  
193 days to decisionK924642 · Product code: **DWF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k924642/>**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 14, 1992
Decision date	Mar 26, 1993
Days to decision	193 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronic Bio-Medicus, Inc.</b>
Location	Eden Prairie, MN, US
Contact	THOMAS K JOHNSEN
510(k) history	13 submissions · 13 cleared · 1992-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k924642/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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