

**K012538 TUBING AND CONNECTORS WITH TRILLIUM
BIOPASSIVE SURFACE**Aug 23, 2001
16 days to decisionK012538 · Product code: **DWF** · Cardiovascular
Source: <https://www.510kdatabase.net/k012538/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Aug 7, 2001
Decision date	Aug 23, 2001
Days to decision	16 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic Vascular
Location	Walker, MI, US
Contact	MARIE HOLM
510(k) history	475 submissions · 453 cleared · 1977-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k012538/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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