

# K103386 MEDTRONIC VASCULAR - HUB MATERIAL MODIFICATION TO THE 5F AND 8F LAUNCHER(R) GUIDE CATHETERS

Feb 11, 2011  
85 days to decisionK103386 · Product code: **DQY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k103386/>

## SUBMISSION DETAILS

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Percutaneous (DQY)
Date received	Nov 18, 2010
Decision date	Feb 11, 2011
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

## APPLICANT

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Company	<b>Medtronic, Inc.</b>
Location	Mounds View, MN, US
Contact	ANU GAUR
Website	<a href="https://www.medtronic.com">https://www.medtronic.com</a>
510(k) history	209 submissions · 208 cleared · 1981-2026

Medtronic, Inc. is a global medical device manufacturer headquartered in Mounds View, United States. The company develops and markets a broad range of medical devices across multiple therapeutic areas. Medtronic maintains a strong FDA 510(k) regulatory track record with FDA 510(k) cleared devices from total submissions since 1981. The company specializes primarily in Cardiovascular devices, which represent 82% of its submission portfolio. Recent clearances include coronary perfusion cannulae, intracoronary shunts, venous cannulae, guidewires, deflectable catheter systems,...