

**K972944 MEDTRONIC MUSTANG SUPER FLOPPY GUIDE  
WIRE/FLOPPY GUIDE WIRE/INTERMEDIATE GUIDE  
WIRE/STANDARD GUIDE WIRE/EXTRA SUPPORT**Nov 3, 1997  
84 days to decisionK972944 · Product code: **DQX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k972944/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Aug 11, 1997
Decision date	Nov 3, 1997
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Medtronics Interventional Vascular</b>
Location	Danvers, MA, US
Contact	MARY DE ARMOND
510(k) history	21 submissions · 21 cleared · 1992-1999

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k972944/> 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