

K082873 MEDTRONIC ANGIOGRAPHIC GUIDE WIRESFeb 6, 2009
130 days to decisionK082873 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k082873/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Sep 29, 2008
Decision date	Feb 6, 2009
Days to decision	130 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k082873/> Data sourced from FDA 510(k) public records (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. - Generated May 13, 2026