

K022764 7F LAUNCHER GUIDE CATHETERAug 28, 2002
7 days to decisionK022764 · Product code: **DQY** · Cardiovascular
Source: <https://www.510kdatabase.net/k022764/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Percutaneous (DQY)
Date received	Aug 21, 2002
Decision date	Aug 28, 2002
Days to decision	7 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k022764/> 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 18, 2026