FDA 510(k) Premarket Notification

K010579 STENT SUPPORT GUIDE CATHETER

CLEARED

Nov 21, 2001
267 days to decision

K010579 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k010579/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Feb 27, 2001
Decision date	Nov 21, 2001
Days to decision	267 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.net
Device record: https://www.510kdatabase.net/k010579/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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