FDA 510(k) Premarket Notification

CLEARED

Jul 31, 2003
150 days to decision

K030671 MEDTRONIOC 7F LAUNCHER BIOPSY GUIDE CATHETER, MODEL LA7RVBB, MEDTRONIC 8F LAUNCHER BIOPSY CATHETER, MODEL LA8RVBB

K030671 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k030671/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Mar 3, 2003
Decision date	Jul 31, 2003
Days to decision	150 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/k030671/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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