FDA 510(k) Premarket Notification

K024069 ONE PIECE PEDIATRIC ARTERIAL CANNULA, MODEL: 77006, 77106

CLEARED

Jan 8, 2003
29 days to decision

K024069 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k024069/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Dec 10, 2002
Decision date	Jan 8, 2003
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medtronic Vascular**
Location	Walker, MI, US
Contact	RONALD W BENNETT
510(k) history	475 submissions · 453 cleared · 1977-2023

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k024069/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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