FDA 510(k) Premarket Notification

K013013 MEDTRONIC DLP ARTERIAL CANNULA WITH 3D TIP - 22 FR.

CLEARED

Dec 4, 2001
88 days to decision

K013013 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k013013/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Sep 7, 2001
Decision date	Dec 4, 2001
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medtronic Cardiac Surgical Products**
Location	Grand Rapids, MI, US
Contact	ROGER W BRINK
510(k) history	7 submissions · 7 cleared · 2000-2003