CLEARED

K010737 SELECT CAP ARTERIAL CANNULA WITH PRESSURE MONITORING LINE, MODEL 78818,78820,78822,78824,78918,78920,78922,78924

Apr 4, 2001
23 days to decision

K010737 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k010737/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Mar 12, 2001
Decision date	Apr 4, 2001
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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