FDA 510(k) Premarket Notification

K980973 MEDTRONIC ANGIOGRAPHIC CATHETER

CLEARED

Dec 10, 1998
269 days to decision

K980973 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k980973/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Mar 16, 1998
Decision date	Dec 10, 1998
Days to decision	269 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medtronics Interventional Vascular**
Location	Danvers, MA, US
Contact	MARK CHARTIER
510(k) history	21 submissions · 21 cleared · 1992-1999