FDA 510(k) Premarket Notification

K925342 MEDTRONIC INTERVENTIONAL VASCULAR SHERPA GUIDEWIRE

CLEARED

May 6, 1993
198 days to decision

K925342 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k925342/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Oct 20, 1992
Decision date	May 6, 1993
Days to decision	198 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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