FDA 510(k) Premarket Notification

K915351 SHERPA GUIDING CATHETER WITH CATHETER INTRODUCER

CLEARED

Jun 29, 1992
216 days to decision

K915351 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k915351/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Nov 26, 1991
Decision date	Jun 29, 1992
Days to decision	216 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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