FDA 510(k) Premarket Notification

K934122 SCIMED TRANSEND EX STEERABLE GUIDE WIRE AND ACCESSORIES

CLEARED

Oct 7, 1993
43 days to decision

K934122 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k934122/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Aug 25, 1993
Decision date	Oct 7, 1993
Days to decision	43 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Scimed Peripheral Interventions**
Location	Plymouth, MN, US
Contact	DARLENE A THOMETZ
510(k) history	8 submissions · 8 cleared · 1993-1995