FDA 510(k) Premarket Notification

K941710 SCIMED TRANSEND STEERABLE GUIDE WIRE AND ACCESSORIES

CLEARED

Sep 1, 1994
148 days to decision

K941710 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k941710/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Apr 6, 1994
Decision date	Sep 1, 1994
Days to decision	148 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