FDA 510(k) Premarket Notification

K992142 SCIMED ANGIOGRAPHIC CATHETERS

CLEARED

Jul 21, 1999
27 days to decision

K992142 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k992142/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jun 24, 1999
Decision date	Jul 21, 1999
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Scimed**
Location	Maple Grove, MN, US
Contact	MELANIE RASKA
510(k) history	8 submissions · 8 cleared · 1990-1999