FDA 510(k) Premarket Notification

K951720 SCIMED EXPO ANGIOGRAPHIC CATHETER

CLEARED

Jun 22, 1995
69 days to decision

K951720 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k951720/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Apr 14, 1995
Decision date	Jun 22, 1995
Days to decision	69 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Scimed Life Systems, Inc.**
Location	Mchenry, IL, US
Contact	DEBORAH L JENSEN
510(k) history	109 submissions · 108 cleared · 1977-1998