FDA 510(k) Premarket Notification

K950094 SCIMED 6 FRENCH MICRO-MAX(TM) GUIDE CATHETER

CLEARED

Mar 3, 1995
52 days to decision

K950094 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k950094/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jan 10, 1995
Decision date	Mar 3, 1995
Days to decision	52 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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