FDA 510(k) Premarket Notification

K980360 SCIMED QUEST FLOPPY AND MODERATE SUPPORT GUIDE WIRES

CLEARED

Apr 29, 1998
90 days to decision

K980360 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k980360/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jan 29, 1998
Decision date	Apr 29, 1998
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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