FDA 510(k) Premarket Notification

K990173 PATHFINDER II ANGIOGRAPHIC CATHETER

CLEARED

Dec 21, 1999
336 days to decision

K990173 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k990173/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jan 19, 1999
Decision date	Dec 21, 1999
Days to decision	336 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Maxxim Medical**
Location	Arlington, TX, US
Contact	EDDIE MONROE
510(k) history	26 submissions · 25 cleared · 1994-2002