FDA 510(k) Premarket Notification

K003902 MULTITRACK ANGIOGRAPHIC CATHETER

CLEARED

Jan 11, 2001
23 days to decision

K003902 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k003902/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Dec 19, 2000
Decision date	Jan 11, 2001
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**NuMED, Inc.**
Location	Hopkinton, NY, US
Contact	NICHELLE R LAFLESH
510(k) history	49 submissions · 47 cleared · 1985-2022