FDA 510(k) Premarket Notification

K013601 NUMED MULLINS PTA CATHETER

CLEARED

Jan 3, 2002
64 days to decision

K013601 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k013601/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Percutaneous (DQY)
Date received	Oct 31, 2001
Decision date	Jan 3, 2002
Days to decision	64 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