

K013601 NUMED MULLINS PTA CATHETERJan 3, 2002
64 days to decisionK013601 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k013601/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 28, 2026