

**K040830 PTA, PTV, ATRIOSEPTOSTOMY, ANGIOGRAPHIC,  
SIZING CATHETERS**Jun 18, 2004  
79 days to decisionK040830 · Product code: **DQO** · Cardiovascular  
Source: <https://www.510kdatabase.net/k040830/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Mar 31, 2004
Decision date	Jun 18, 2004
Days to decision	79 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k040830/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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