

**K790083 CATHETER, ARTERIAL LINE**Jan 25, 1979  
13 days to decisionK790083 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k790083/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jan 12, 1979
Decision date	Jan 25, 1979
Days to decision	13 days
Third-party review	No

**APPLICANT**

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Company	<b>Namic</b>
Location	Walker, MI, US
510(k) history	8 submissions · 8 cleared · 1979-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k790083/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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