

**K884375 TRANSLUMBAR AORTOGRAPHY NEEDLE CATHETER**Jan 11, 1989  
85 days to decisionK884375 · Product code: **DQR** · CardiovascularSource: <https://www.510kdatabase.net/k884375/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Catheter (DQR)
Date received	Oct 18, 1988
Decision date	Jan 11, 1989
Days to decision	85 days
Third-party review	No

**APPLICANT**

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Company	<b>Argon Medical Corp.</b>
Location	Mchenry, IL, US
Contact	BRISTER, P.E.
510(k) history	27 submissions · 27 cleared · 1976-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k884375/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 29, 2026