

**K882221 OVER THE NEEDLE CATHETER, ARTERIAL  
CATHETER NEEDLE**Aug 12, 1988  
78 days to decisionK882221 · Product code: **DYB** · Cardiovascular  
Source: <https://www.510kdatabase.net/k882221/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	May 26, 1988
Decision date	Aug 12, 1988
Days to decision	78 days
Third-party review	No

**APPLICANT**

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Company	<b>Argon Medical Corp.</b>
Location	Mchenry, IL, US
Contact	DAVID MEYERS
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/k882221/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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