

**K875353 PERCUTANEOUS ARTERIAL/VENOUS CANNULA  
PLACEMENT KIT**Mar 22, 1988  
82 days to decisionK875353 · Product code: **DWF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k875353/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Dec 31, 1987
Decision date	Mar 22, 1988
Days to decision	82 days
Third-party review	No

**APPLICANT**

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Company	<b>Dip, Inc.</b>
Location	Mchenry, IL, US
Contact	VAN HOF
510(k) history	56 submissions · 56 cleared · 1979-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k875353/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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