

**K874852 POLYSTAN WIRE REINFORCED TWO STAGE  
CATHETERS**Feb 5, 1988  
72 days to decisionK874852 · Product code: **DWF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k874852/>**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	Nov 25, 1987
Decision date	Feb 5, 1988
Days to decision	72 days
Third-party review	No

**APPLICANT**

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Company	<b>Polystan C/O Vitalcor, Inc.</b>
Location	Ballerup Denmark, US
Contact	BILL HUCK
510(k) history	8 submissions · 8 cleared · 1984-1990

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k874852/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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