

**K001509 STOCKERT V172 SERIES VENOUS FEMORAL CANNULAE**Oct 26, 2001  
529 days to decisionK001509 · Product code: **DWF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k001509/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	May 15, 2000
Decision date	Oct 26, 2001
Days to decision	529 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cobe Cardiovascular, Inc.</b>
Location	Arvada, CO, US
Contact	LYNNE LEONARD
510(k) history	43 submissions · 43 cleared · 1992-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k001509/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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