

**K964194 CHASE CURVED LEFT HEART VENT CATHETER**Apr 16, 1997  
177 days to decisionK964194 · Product code: **DWF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k964194/>**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	Oct 21, 1996
Decision date	Apr 16, 1997
Days to decision	177 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Chase Medical, Inc.</b>
Location	Richardson, TX, US
Contact	BERT DAVIS
510(k) history	27 submissions · 27 cleared · 1997-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k964194/> 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 May 19, 2026