

**K964198 CAHSE AORTIC ROOT CANNULA**Mar 19, 1997  
149 days to decisionK964198 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k964198/>**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	Mar 19, 1997
Days to decision	149 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/k964198/> 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