

**K981995 QUICKDRAW VENOUS CANNULA**Apr 20, 1999  
316 days to decisionK981995 · Product code: **DWF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k981995/>**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	Jun 8, 1998
Decision date	Apr 20, 1999
Days to decision	316 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Heartport, Inc.</b>
Location	Redwood City, CA, US
Contact	MARIANNE C DRENNAN
510(k) history	24 submissions · 24 cleared · 1996-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k981995/> 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 July 4, 2026