

K961245 HEARTPORT ENDOPULMONARY VENTJun 18, 1996
78 days to decisionK961245 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k961245/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Apr 1, 1996
Decision date	Jun 18, 1996
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Heartport, Inc.
Location	Redwood City, CA, US
Contact	ROBERT J CHIN
510(k) history	24 submissions · 24 cleared · 1996-2000

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