

K962835 HEARTPORT ENDOVENOUS DRAINAGE CANNULAFeb 10, 1997
203 days to decisionK962835 · Product code: **DWF** · Cardiovascular
Source: <https://www.510kdatabase.net/k962835/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Jul 22, 1996
Decision date	Feb 10, 1997
Days to decision	203 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k962835/> Data sourced from FDA 510(k) public records (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. - Generated July 4, 2026