

K831332 ARGYLE DUAL STAGE VENOUS RETURN CATH.Jun 30, 1983
66 days to decisionK831332 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k831332/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Apr 25, 1983
Decision date	Jun 30, 1983
Days to decision	66 days
Third-party review	No

APPLICANT

Company	Sherwood Medical Co.
Location	Mchenry, IL, US
510(k) history	191 submissions · 177 cleared · 1976-1998

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Device record: <https://www.510kdatabase.net/k831332/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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