FDA 510(k) Premarket Notification

K831332 ARGYLE DUAL STAGE VENOUS RETURN CATH.

CLEARED

Jun 30, 1983
66 days to decision

K831332 · Product code: **DWF** · Cardiovascular
Source: 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