

K844137 STOCKERT-SHILEY VENOUS CATHETER & AORTIC ARCH CANNDec 27, 1984
65 days to decisionK844137 · Product code: **DWF** · Cardiovascular
Source: <https://www.510kdatabase.net/k844137/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Oct 23, 1984
Decision date	Dec 27, 1984
Days to decision	65 days
Third-party review	No

APPLICANT

Company	Shiley, Inc.
Location	Mchenry, IL, US
Contact	ROBERT CURTIS
510(k) history	174 submissions · 174 cleared · 1976-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k844137/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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