FDA 510(k) Premarket Notification

K844137 STOCKERT-SHILEY VENOUS CATHETER & AORTIC ARCH CANN

CLEARED

Dec 27, 1984
65 days to decision

K844137 · 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