FDA 510(k) Premarket Notification

K881074 SHILEY FEMORAL VENOUS CANNULA

CLEARED

May 31, 1988
78 days to decision

K881074 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k881074/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Mar 14, 1988
Decision date	May 31, 1988
Days to decision	78 days
Third-party review	No

APPLICANT

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