FDA 510(k) Premarket Notification

K920902 MODIFICATION OF SARNS TWO STAGE VENOUS CANNULA

CLEARED

May 11, 1993
440 days to decision

K920902 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k920902/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Feb 26, 1992
Decision date	May 11, 1993
Days to decision	440 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**3M Health Care, Sarns**
Location	Mchenry, IL, US
Contact	CATHY L SIMPSON
510(k) history	76 submissions · 76 cleared · 1976-1996