FDA 510(k) Premarket Notification

K915193 SARNS RETROGRADE CANNULA

CLEARED

Mar 4, 1992
106 days to decision

K915193 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k915193/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Nov 19, 1991
Decision date	Mar 4, 1992
Days to decision	106 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