

K915193 SARNS RETROGRADE CANNULAMar 4, 1992
106 days to decisionK915193 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k915193/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 6, 2026