

K770431 CATHETER, TWO STAGE VENOUS RETURNMar 16, 1977
9 days to decisionK770431 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k770431/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Mar 7, 1977
Decision date	Mar 16, 1977
Days to decision	9 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k770431/> 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 1, 2026