

K854640 MODIFIED CORONARY CARDIOPLEGIA ADAPTORFeb 10, 1986
83 days to decisionK854640 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k854640/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Nov 19, 1985
Decision date	Feb 10, 1986
Days to decision	83 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k854640/>, 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 May 26, 2026