

K831591 AORTIC ROOT CANNULA W/INTEGRAL PRESSJun 30, 1983
44 days to decisionK831591 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k831591/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	May 17, 1983
Decision date	Jun 30, 1983
Days to decision	44 days
Third-party review	No

APPLICANT

Company	Dlp, Inc.
Location	Mchenry, IL, US
510(k) history	56 submissions · 56 cleared · 1979-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k831591/> 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 2, 2026