

K821149 SUBMISSION CORONARY CANNULAMay 18, 1982
25 days to decisionK821149 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k821149/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Apr 23, 1982
Decision date	May 18, 1982
Days to decision	25 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k821149/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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