

K810548 CARDIOPULMINARY BYPASS CATHETER CANNULAMar 17, 1981
15 days to decisionK810548 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k810548/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Mar 2, 1981
Decision date	Mar 17, 1981
Days to decision	15 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/k810548/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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