

K790565 AORTIC ROOT CANNULAJun 11, 1979
84 days to decisionK790565 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k790565/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Mar 19, 1979
Decision date	Jun 11, 1979
Days to decision	84 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/k790565/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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