

**K810820 CARDIOPULMONARY BYPASS**Apr 3, 1981  
10 days to decisionK810820 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k810820/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Mar 24, 1981
Decision date	Apr 3, 1981
Days to decision	10 days
Third-party review	No

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k810820/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 2, 2026