FDA 510(k) Premarket Notification

K810820 CARDIOPULMONARY BYPASS

CLEARED

Apr 3, 1981
10 days to decision

K810820 · Product code: **DWF** · Cardiovascular
Source: 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	**Dlp, Inc.**
Location	Mchenry, IL, US
510(k) history	56 submissions · 56 cleared · 1979-1997