FDA 510(k) Premarket Notification

K875353 PERCUTANEOUS ARTERIAL/VENOUS CANNULA PLACEMENT KIT

CLEARED

Mar 22, 1988
82 days to decision

K875353 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k875353/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Dec 31, 1987
Decision date	Mar 22, 1988
Days to decision	82 days
Third-party review	No

APPLICANT

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