FDA 510(k) Premarket Notification

K810515 DISPOSABLE 2-STAGE CANNULA

CLEARED

Apr 23, 1981
57 days to decision

K810515 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k810515/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Feb 25, 1981
Decision date	Apr 23, 1981
Days to decision	57 days
Third-party review	No

APPLICANT

Company	**Texas Medical Products, Inc.**
Location	Mchenry, IL, US
510(k) history	30 submissions · 30 cleared · 1980-1989