FDA 510(k) Premarket Notification

CLEARED

K001509 STOCKERT V172 SERIES VENOUS FEMORAL CANNULAE

Oct 26, 2001
529 days to decision

K001509 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k001509/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	May 15, 2000
Decision date	Oct 26, 2001
Days to decision	529 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cobe Cardiovascular, Inc.**
Location	Arvada, CO, US
Contact	LYNNE LEONARD
510(k) history	43 submissions · 43 cleared · 1992-2005