FDA 510(k) Premarket Notification

K994209 STOCKERT V142 SERIES VENOUS CANNULAE WITH LIGHTHOUSE TIP

CLEARED

Jun 14, 2000
183 days to decision

K994209 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k994209/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Dec 14, 1999
Decision date	Jun 14, 2000
Days to decision	183 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