FDA 510(k) Premarket Notification

K964198 CAHSE AORTIC ROOT CANNULA

CLEARED

Mar 19, 1997
149 days to decision

K964198 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k964198/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Oct 21, 1996
Decision date	Mar 19, 1997
Days to decision	149 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Chase Medical, Inc.**
Location	Richardson, TX, US
Contact	BERT DAVIS
510(k) history	27 submissions · 27 cleared · 1997-2004