FDA 510(k) Premarket Notification

K964194 CHASE CURVED LEFT HEART VENT CATHETER

CLEARED

Apr 16, 1997
177 days to decision

K964194 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k964194/

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	Apr 16, 1997
Days to decision	177 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