FDA 510(k) Premarket Notification

K992932 CIPA (ADULTS) CIPI (INFANTS)

CLEARED

Jun 7, 2000
281 days to decision

K992932 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k992932/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Aug 31, 1999
Decision date	Jun 7, 2000
Days to decision	281 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Dmc Medical, Ltd.**
Location	Fullerton, CA, US
Contact	CHARMAINE HENDERSON
510(k) history	3 submissions · 3 cleared · 2000-2011