

K992932 CIPA (ADULTS) CIPI (INFANTS)Jun 7, 2000
281 days to decisionK992932 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k992932/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 19, 2026