

K001371 CARDEON ASCENDING BALLOON CANNULA (ABC)Jul 7, 2000
67 days to decisionK001371 · Product code: **DWF** · Cardiovascular
Source: <https://www.510kdatabase.net/k001371/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	May 1, 2000
Decision date	Jul 7, 2000
Days to decision	67 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cardeon Corp.
Location	Cupertino, CA, US
Contact	JANE BEGGS
510(k) history	6 submissions · 6 cleared · 2000-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k001371/> 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 July 3, 2026