

K042156 CARDEON COBRA CATHETEROct 6, 2004
57 days to decisionK042156 · Product code: **DWF** · Cardiovascular
Source: <https://www.510kdatabase.net/k042156/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Aug 10, 2004
Decision date	Oct 6, 2004
Days to decision	57 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/k042156/> 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