FDA 510(k) Premarket Notification

K022174 MODIFICATION TO CARDEON AEGIS CATHETER

CLEARED

Jul 29, 2002
26 days to decision

K022174 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k022174/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Jul 3, 2002
Decision date	Jul 29, 2002
Days to decision	26 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.net
Device record: https://www.510kdatabase.net/k022174/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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