

**K992965 CORDIS WEBSTER FIXED CURVE CATHETERS,
MODELS D-1124, D-1085**Nov 26, 1999
85 days to decisionK992965 · Product code: DRF · Cardiovascular
Source: <https://www.510kdatabase.net/k992965/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Sep 2, 1999
Decision date	Nov 26, 1999
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cordis Webster, Inc.
Location	Baldwin Park, CA, US
Contact	MARY ADAMS
510(k) history	10 submissions · 10 cleared · 1995-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k992965/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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