

**K982415 CORDIS WEBSTER REF-STAR EXTERNAL
REFERENCE PATCH, MODEL # D-1210**Aug 10, 1998
28 days to decisionK982415 · Product code: DRF · Cardiovascular
Source: <https://www.510kdatabase.net/k982415/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Jul 13, 1998
Decision date	Aug 10, 1998
Days to decision	28 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/k982415/> 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 4, 2026