

**K955817 CORIDIS WEBSTER DEFLECTABLE BRAIDED-TIP  
ELECTRODE CATHETER**Mar 25, 1996  
90 days to decisionK955817 · Product code: **DRF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k955817/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Dec 26, 1995
Decision date	Mar 25, 1996
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k955817/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated July 4, 2026