

**K991531 CORDIS WEBSTER DIAGNOSTIC DEFLECTABLE TIP
CATHETER, MODEL D-1078**Oct 4, 1999
154 days to decisionK991531 · Product code: DRF · Cardiovascular
Source: <https://www.510kdatabase.net/k991531/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	May 3, 1999
Decision date	Oct 4, 1999
Days to decision	154 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/k991531/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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