

**K953768 CORDIS WEBSTER A20 DIAGNOSTIC DEFLECTABLE
TIP CATHETER**Nov 22, 1995
100 days to decisionK953768 · Product code: **DRF** · Cardiovascular
Source: <https://www.510kdatabase.net/k953768/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Aug 14, 1995
Decision date	Nov 22, 1995
Days to decision	100 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/k953768/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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