

K953663 CORDIS WEBSTER T20 DIAGNOSTIC DEFLECTABLE TIP CATHETERSep 18, 1996
408 days to decisionK953663 · Product code: **DRF** · Cardiovascular
Source: <https://www.510kdatabase.net/k953663/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Aug 7, 1995
Decision date	Sep 18, 1996
Days to decision	408 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/k953663/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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