FDA 510(k) Premarket Notification

K955817 CORIDIS WEBSTER DEFLECTABLE BRAIDED-TIP ELECTRODE CATHETER

CLEARED

Mar 25, 1996
90 days to decision

K955817 · Product code: **DRF** · Cardiovascular
Source: https://www.510kdatabase.net/k955817/

SUBMISSION DETAILS

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

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