FDA 510(k) Premarket Notification

**CLEARED**

# K992965 CORDIS WEBSTER FIXED CURVE CATHETERS, MODELS D-1124, D-1085

Nov 26, 1999
85 days to decision

K992965 · Product code: **DRF** · Cardiovascular
Source: https://www.510kdatabase.net/k992965/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF) |
| Date received | Sep 2, 1999 |
| Decision date | Nov 26, 1999 |
| Days to decision | 85 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 |