FDA 510(k) Premarket Notification

# K882593 MODIFIED CONTINUOUS CARDIAC OUTPUT MONITOR SYSTEM

**CLEARED**

Aug 29, 1988
66 days to decision

K882593 · Product code: **DPW** · Cardiovascular
Source: https://www.510kdatabase.net/k882593/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Flowmeter, Blood, Cardiovascular (DPW) |
| Date received | Jun 24, 1988 |
| Decision date | Aug 29, 1988 |
| Days to decision | 66 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Cardio Metrics, Inc.** |
| Location | Houston, TX, US |
| Contact | MICHAEL J BILLIG |
| 510(k) history | 23 submissions · 23 cleared · 1986-1997 |