

**K882593 MODIFIED CONTINUOUS CARDIAC OUTPUT  
MONITOR SYSTEM**Aug 29, 1988  
66 days to decisionK882593 · 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	<b>Cardio Metrics, Inc.</b>
Location	Houston, TX, US
Contact	MICHAEL J BILLIG
510(k) history	23 submissions · 23 cleared · 1986-1997

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882593/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 20, 2026