

**K901733 MODIFIED FLOCATH PULMONARY ARTERY
CATHETER**Jul 16, 1990
90 days to decisionK901733 · Product code: **DPW** · Cardiovascular
Source: <https://www.510kdatabase.net/k901733/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Flowmeter, Blood, Cardiovascular (DPW)
Date received	Apr 17, 1990
Decision date	Jul 16, 1990
Days to decision	90 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k901733/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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