

**K912446 MODIF. LASERFLO BLOOD PERFUSION MONITOR
MODEL BPM2**

Dec 12, 1991
192 days to decision

K912446 · Product code: **DPW** · Cardiovascular
Source: <https://www.510kdatabase.net/k912446/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Flowmeter, Blood, Cardiovascular (DPW)
Date received	Jun 3, 1991
Decision date	Dec 12, 1991
Days to decision	192 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Vasamedics, LLC
Location	Minneapolis, MN, US
Contact	CONSTANCE G BUNDY
510(k) history	6 submissions · 5 cleared · 1991-1997

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k912446/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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