

**K911763 LASERFLO(R) BLOOD PERFUSION MONITOR (MODEL BPM2)**

Aug 13, 1991  
116 days to decision

K911763 · Product code: **DPW** · Cardiovascular  
Source: <https://www.510kdatabase.net/k911763/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Flowmeter, Blood, Cardiovascular (DPW)
Date received	Apr 19, 1991
Decision date	Aug 13, 1991
Days to decision	116 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k911763/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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