FDA 510(k) Premarket Notification

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

CLEARED

Aug 13, 1991
116 days to decision

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

SUBMISSION DETAILS

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

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