

K840522 RECOMED BLOOD-FLOW METER H236-052-02May 16, 1985
464 days to decisionK840522 · Product code: **DPW** · CardiovascularSource: <https://www.510kdatabase.net/k840522/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Flowmeter, Blood, Cardiovascular (DPW)
Date received	Feb 7, 1984
Decision date	May 16, 1985
Days to decision	464 days
Third-party review	No

APPLICANT

Company	Litton Medical Electronics
Location	Walker, MI, US
Contact	ROBERT L CASARSA
510(k) history	38 submissions · 38 cleared · 1982-1985

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k840522/>; 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 27, 2026