FDA 510(k) Premarket Notification

K840522 RECOMED BLOOD-FLOW METER H236-052-02

CLEARED

May 16, 1985
464 days to decision

K840522 · Product code: **DPW** · Cardiovascular
Source: 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