FDA 510(k) Premarket Notification

K823404 VOLUMERIC CONTROLLER

CLEARED

Dec 22, 1982
37 days to decision

K823404 · Product code: **LDR** · General Hospital
Source: https://www.510kdatabase.net/k823404/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	Nov 15, 1982
Decision date	Dec 22, 1982
Days to decision	37 days
Third-party review	No

APPLICANT

Company	**Quest Medical, Inc.**
Location	Walker, MI, US
510(k) history	39 submissions · 39 cleared · 1980-2024

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k823404/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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