FDA 510(k) Premarket Notification

K832702 POCKET AID

CLEARED

Nov 17, 1983
98 days to decision

K832702 · Product code: **LDR** · General Hospital
Source: https://www.510kdatabase.net/k832702/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	Aug 11, 1983
Decision date	Nov 17, 1983
Days to decision	98 days
Third-party review	No

APPLICANT

Company	**M.C. Johnson Co., Inc.**
Location	Walker, MI, US
510(k) history	3 submissions · 3 cleared · 1983-1995

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k832702/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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