FDA 510(k) Premarket Notification

K834528 MODEL 2001 INTELLIGENT INFUSOR

CLEARED

Feb 21, 1984
62 days to decision

K834528 · Product code: **LDR** · General Hospital
Source: https://www.510kdatabase.net/k834528/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	Dec 21, 1983
Decision date	Feb 21, 1984
Days to decision	62 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/k834528/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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