FDA 510(k) Premarket Notification

K843254 DK*300 VOLUMETRIC INFUSION CONTROLLER

CLEARED

Oct 11, 1984
52 days to decision

K843254 · Product code: **LDR** · General Hospital
Source: https://www.510kdatabase.net/k843254/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	Aug 20, 1984
Decision date	Oct 11, 1984
Days to decision	52 days
Third-party review	No

APPLICANT

Company	**Deka Research & Development Corp.**
Location	Mchenry, IL, US
Contact	STEPHEN W SAGON
510(k) history	10 submissions · 10 cleared · 1984-2025

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k843254/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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