

K242693 DEKA Infusion System, DEKA Administration SetOct 7, 2024
28 days to decisionK242693 · Product code: **LDR** · General Hospital
Source: <https://www.510kdatabase.net/k242693/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	Sep 9, 2024
Decision date	Oct 7, 2024
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Deka Research and Development Corporation
Location	Manchester, NH, US
Contact	Paul Smolenski
510(k) history	1 submissions · 1 cleared · 2024-2024

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