FDA 510(k) Premarket Notification

K242693 DEKA Infusion System, DEKA Administration Set

CLEARED

Oct 7, 2024
28 days to decision

K242693 · 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