FDA 510(k) Premarket Notification

K150687 DripAssist

CLEARED

Oct 13, 2015
210 days to decision

K150687 · Product code: **FLN** · General Hospital
Source: https://www.510kdatabase.net/k150687/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Electric For Gravity Flow Infusion Systems (FLN)
Date received	Mar 17, 2015
Decision date	Oct 13, 2015
Days to decision	210 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Shift Labs**
Location	Seattle, WA, US
Contact	Beth Kolko
510(k) history	1 submissions · 1 cleared · 2015-2015