FDA 510(k) Premarket Notification

K821823 I.V. SENTINEL

CLEARED

Aug 16, 1982
55 days to decision

K821823 · Product code: **FLN** · General Hospital
Source: https://www.510kdatabase.net/k821823/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Electric For Gravity Flow Infusion Systems (FLN)
Date received	Jun 22, 1982
Decision date	Aug 16, 1982
Days to decision	55 days
Third-party review	No

APPLICANT

Company	**The John Bunn Co.**
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1976-1990