

**K821823 I.V. SENTINEL**Aug 16, 1982  
55 days to decisionK821823 · Product code: **FLN** · General HospitalSource: <https://www.510kdatabase.net/k821823/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k821823/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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