

**K821624 LOW VOLUME INFUSTION SET**Jun 22, 1982  
20 days to decisionK821624 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k821624/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jun 2, 1982
Decision date	Jun 22, 1982
Days to decision	20 days
Third-party review	No

**APPLICANT**

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Company	<b>Lifemed of California</b>
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1982-2001

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

Device record: <https://www.510kdatabase.net/k821624/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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