

**K831315 TRILEX**Aug 12, 1983  
112 days to decisionK831315 · Product code: **FPB** · General Hospital  
Source: <https://www.510kdatabase.net/k831315/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filter, Infusion Line (FPB)
Date received	Apr 22, 1983
Decision date	Aug 12, 1983
Days to decision	112 days
Third-party review	No

**APPLICANT**

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Company	<b>Tri-Med, Inc.</b>
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1977-2004

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

Device record: <https://www.510kdatabase.net/k831315/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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