

**K823690 IVFLO-HP & IVFIL-HP**Dec 28, 1982  
20 days to decisionK823690 · Product code: **FPB** · General HospitalSource: <https://www.510kdatabase.net/k823690/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filter, Infusion Line (FPB)
Date received	Dec 8, 1982
Decision date	Dec 28, 1982
Days to decision	20 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/k823690/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 20, 2026