

K813001 FIBRI-FILTER 20Nov 6, 1981
11 days to decisionK813001 · Product code: **FPB** · General Hospital
Source: <https://www.510kdatabase.net/k813001/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filter, Infusion Line (FPB)
Date received	Oct 26, 1981
Decision date	Nov 6, 1981
Days to decision	11 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k813001/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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