

K831350 IVFLO-HP & IVFIL-HP FILTERAug 12, 1983
109 days to decisionK831350 · Product code: **FPB** · General HospitalSource: <https://www.510kdatabase.net/k831350/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filter, Infusion Line (FPB)
Date received	Apr 25, 1983
Decision date	Aug 12, 1983
Days to decision	109 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k831350/> Data sourced from FDA 510(k) public records (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. - Generated May 20, 2026