

K842075 CRITIKON I.V. FILTERJun 28, 1984
36 days to decisionK842075 · Product code: **FPB** · General Hospital
Source: <https://www.510kdatabase.net/k842075/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filter, Infusion Line (FPB)
Date received	May 23, 1984
Decision date	Jun 28, 1984
Days to decision	36 days
Third-party review	No

APPLICANT

Company	Critikon Company, LLC
Location	Mchenry, IL, US
510(k) history	51 submissions · 51 cleared · 1979-2000

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Device record: <https://www.510kdatabase.net/k842075/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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