

K781785 I.V. SET WITH IN-LINE FILTERDec 7, 1978
50 days to decisionK781785 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k781785/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Oct 18, 1978
Decision date	Dec 7, 1978
Days to decision	50 days
Third-party review	No

APPLICANT

Company	Cutler Laboratories, Inc.
Location	MD, US
510(k) history	2 submissions · 2 cleared · 1978-1978

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Device record: <https://www.510kdatabase.net/k781785/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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