

K791529 INFUSION NEEDLE SETOct 2, 1979
53 days to decisionK791529 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k791529/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 10, 1979
Decision date	Oct 2, 1979
Days to decision	53 days
Third-party review	No

APPLICANT

Company	I.M.P. Group , Ltd.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1979-1979

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Device record: <https://www.510kdatabase.net/k791529/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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