

K811107 IV SETSJun 12, 1981
50 days to decisionK811107 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k811107/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 23, 1981
Decision date	Jun 12, 1981
Days to decision	50 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/k811107/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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