

K810721 PROCESS CHANGEApr 23, 1981
37 days to decisionK810721 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k810721/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 17, 1981
Decision date	Apr 23, 1981
Days to decision	37 days
Third-party review	No

APPLICANT

Company	Medlon, Inc.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981

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Device record: <https://www.510kdatabase.net/k810721/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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