

**K801821 FLOW MED 150**Aug 13, 1980  
13 days to decisionK801821 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k801821/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jul 31, 1980
Decision date	Aug 13, 1980
Days to decision	13 days
Third-party review	No

**APPLICANT**

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Company	<b>D.R.O. Medical, Inc.</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1980-1985

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k801821/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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