

**K801004 FLOW MED 160**May 2, 1980  
7 days to decisionK801004 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k801004/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 25, 1980
Decision date	May 2, 1980
Days to decision	7 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/k801004/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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