

**K781637 MODEL IV 1500 STANDARD BURETTE**Dec 12, 1978  
82 days to decisionK781637 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k781637/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 21, 1978
Decision date	Dec 12, 1978
Days to decision	82 days
Third-party review	No

**APPLICANT**

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Company	<b>Valleylab, Inc.</b>
Location	Mchenry, IL, US
510(k) history	94 submissions · 93 cleared · 1976-2003

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

Device record: <https://www.510kdatabase.net/k781637/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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