

**K833178 SOLUTION ADMINISTRATION LINE W/SPIKE**Nov 30, 1983  
72 days to decisionK833178 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k833178/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 19, 1983
Decision date	Nov 30, 1983
Days to decision	72 days
Third-party review	No

**APPLICANT**

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Company	<b>Intramed, Inc.</b>
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1983-1983

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Device record: <https://www.510kdatabase.net/k833178/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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