

**K832542 I.V. ADMINISTRATION SET**Oct 31, 1983  
94 days to decisionK832542 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k832542/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jul 29, 1983
Decision date	Oct 31, 1983
Days to decision	94 days
Third-party review	No

**APPLICANT**

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Company	<b>Research Industries Corp.</b>
Location	Walker, MI, US
510(k) history	5 submissions · 5 cleared · 1983-1995

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

Device record: <https://www.510kdatabase.net/k832542/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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