

**K901949 IV SET**Sep 12, 1990  
135 days to decisionK901949 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k901949/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 30, 1990
Decision date	Sep 12, 1990
Days to decision	135 days
Third-party review	No

**APPLICANT**

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Company	<b>Jfv, Inc.</b>
Location	Spencer, IN, US
Contact	VANCE, SR.
510(k) history	1 submissions · 1 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k901949/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 7, 2026