

**K904013 SPIVE NEEDLE SYSTEM, MODIFICATION**Oct 1, 1990  
33 days to decisionK904013 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k904013/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 29, 1990
Decision date	Oct 1, 1990
Days to decision	33 days
Third-party review	No

**APPLICANT**

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Company	<b>Pascall Medical Corp.</b>
Location	Bethesda, MD, US
Contact	WILLIAM R TARELLO
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/k904013/>; 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