

**K905219 &apos; LOW VOLUME IV EXTENSION SET**Jun 4, 1991  
196 days to decisionK905219 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k905219/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Nov 20, 1990
Decision date	Jun 4, 1991
Days to decision	196 days
Third-party review	No

**APPLICANT**

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Company	<b>Marquette Medical, Inc.</b>
Location	Crofton, MD, US
Contact	MARQUETTE JR
510(k) history	17 submissions · 17 cleared · 1987-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k905219/>; 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 1, 2026