

**K965030 BRANNON ARTERIO-VEINUS PORTSYRINGE**Mar 4, 1997  
77 days to decisionK965030 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k965030/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Dec 17, 1996
Decision date	Mar 4, 1997
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vascular Logics, Inc.</b>
Location	Huntington Beach, CA, US
Contact	JAMES K BRANNON
510(k) history	1 submissions · 1 cleared · 1997-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k965030/> 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 6, 2026