

**K052075 PROCEDUR-10**Nov 9, 2005  
100 days to decisionK052075 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k052075/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Syringe, Piston (FMF)
Date received	Aug 1, 2005
Decision date	Nov 9, 2005
Days to decision	100 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Avanca Medical Devices, Inc.</b>
Location	Albuquerque, NM, US
Contact	FRANK FERGUSON
510(k) history	3 submissions · 3 cleared · 2005-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k052075/> 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 May 21, 2026