

**K945592 VLV SOLODRAW(TM) DUAL CHAMBER PISTON  
SYRINGE**Jul 14, 1995  
242 days to decisionK945592 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k945592/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Nov 14, 1994
Decision date	Jul 14, 1995
Days to decision	242 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Vlv Associates, Inc.</b>
Location	East Hanover, NJ, US
Contact	VINCENT L VAILANCOURT
510(k) history	12 submissions · 11 cleared · 1986-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k945592/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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