

**K043261 HARVEST GRAFT DELIVERY SYRINGE**Mar 11, 2005  
107 days to decisionK043261 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k043261/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Nov 24, 2004
Decision date	Mar 11, 2005
Days to decision	107 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Harvest Technologies, Corp.</b>
Location	Plymouth, MA, US
Contact	JACK BONASERA
510(k) history	6 submissions · 6 cleared · 2001-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k043261/> 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 June 13, 2026