

**K020252 SMARTJET GRAFTING LIQUID APPLICATOR,  
MODELS SK/S & LK/2**Apr 5, 2002  
71 days to decisionK020252 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k020252/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jan 24, 2002
Decision date	Apr 5, 2002
Days to decision	71 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Harvest Technologies, Corp.</b>
Location	Plymouth, MA, US
Contact	JOHN D 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/k020252/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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