

**K090929 STERILE HYPODERMIC SYRINGE AND NEEDLE FOR SINGLE USE, STERILE INSULING SYRINGE FOR SINGLE USE WITH FIXED NEEDLE**

Jun 12, 2009  
71 days to decision

K090929 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k090929/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Apr 2, 2009
Decision date	Jun 12, 2009
Days to decision	71 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Shanghai Double Dove Industrial Co., Ltd.</b>
Location	Shanghai, CN
Contact	Diana Hong
510(k) history	2 submissions · 2 cleared · 2009-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k090929/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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