

**K113673 MYSAFETY SYRINGE**Aug 3, 2012  
234 days to decisionK113673 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k113673/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Dec 13, 2011
Decision date	Aug 3, 2012
Days to decision	234 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medicalchain Internartional Corporation</b>
Location	Thousand Oaks, CA, US
Contact	ROBIN HWANG
510(k) history	1 submissions · 1 cleared · 2012-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k113673/> 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 14, 2026