

**K101708 PATIENT SAFE LUER CAP**Feb 24, 2011  
252 days to decisionK101708 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k101708/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jun 17, 2010
Decision date	Feb 24, 2011
Days to decision	252 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Retractable Technologies, Inc.</b>
Location	Lewisville, TX, US
Contact	RHONDA WELLS
510(k) history	13 submissions · 13 cleared · 1995-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k101708/> 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 May 27, 2026