

**K122416 ARTIGLASS L. O.R. GLASS SYRINGE**May 3, 2013  
268 days to decisionK122416 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k122416/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Aug 8, 2012
Decision date	May 3, 2013
Days to decision	268 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Artiglass Srl</b>
Location	Due Carrare Padova, IT
Contact	FEDERICO BACCARIN
510(k) history	2 submissions · 2 cleared · 2013-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k122416/> 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 26, 2026