

**K201338 Allograft MIS Delivery System**Jul 13, 2020  
54 days to decisionK201338 · Product code: **FMF** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k201338/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	May 20, 2020
Decision date	Jul 13, 2020
Days to decision	54 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Lifenet Health</b>
Location	Virginia Beach, VA, US
Contact	Brittany Beasley
510(k) history	3 submissions · 3 cleared · 2008-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k201338/> 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