

**K833460 SYRINGE,PISTON NEEDLE, HOLLOW LUMEN**Nov 28, 1983  
54 days to decisionK833460 · Product code: **FMF** · General HospitalSource: <https://www.510kdatabase.net/k833460/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Oct 5, 1983
Decision date	Nov 28, 1983
Days to decision	54 days
Third-party review	No

**APPLICANT**

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Company	<b>Aldrich Chemical Co., Inc.</b>
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1983-1983

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k833460/> 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 July 9, 2026