

**K895203 EN-TRAP SYRINGE**Oct 18, 1989  
61 days to decisionK895203 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k895203/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Aug 18, 1989
Decision date	Oct 18, 1989
Days to decision	61 days
Third-party review	No

**APPLICANT**

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Company	<b>The Medtech Group, Inc.</b>
Location	South Plainfield, NJ, US
Contact	LAURENCE G WESSER
510(k) history	2 submissions · 2 cleared · 1986-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895203/>; 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