

**K923321 EDGE SYRINGES AND NEEDLES**Sep 28, 1993  
448 days to decisionK923321 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k923321/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jul 7, 1992
Decision date	Sep 28, 1993
Days to decision	448 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Trico Intl., Inc.</b>
Location	New York, NY, US
Contact	ESTHER LAI
510(k) history	2 submissions · 2 cleared · 1992-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k923321/> 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 7, 2026