

**K923127 NEEDLE-PRO CARTRIDGE**Apr 29, 1993  
307 days to decisionK923127 · Product code: **FMI** · General Hospital  
Source: <https://www.510kdatabase.net/k923127/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jun 26, 1992
Decision date	Apr 29, 1993
Days to decision	307 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Concord/Portex</b>
Location	Keene, NH, US
Contact	ROBERT WHEELER
510(k) history	23 submissions · 20 cleared · 1989-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k923127/> 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