

**K960698 NON-CORING NEEDLE**Apr 15, 1996  
55 days to decisionK960698 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k960698/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Feb 20, 1996
Decision date	Apr 15, 1996
Days to decision	55 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Med Institute, Inc.</b>
Location	West Lafayette, IN, US
Contact	NEAL E FEARNOT, PH.D., E.E.
510(k) history	26 submissions · 23 cleared · 1990-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960698/> 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 20, 2026