

**K980003 M500-100 SHARPS CONTAINER**Mar 13, 1998  
70 days to decisionK980003 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k980003/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jan 2, 1998
Decision date	Mar 13, 1998
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Imagination Medical, Inc.</b>
Location	Jacksonville, FL, US
Contact	TIM WEIST
510(k) history	1 submissions · 1 cleared · 1998-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k980003/> 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 5, 2026