

**K992404 THE WAND**Jun 6, 2000  
323 days to decisionK992404 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k992404/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jul 19, 1999
Decision date	Jun 6, 2000
Days to decision	323 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Spintech, Inc.</b>
Location	Minneapolis, MN, US
Contact	DANIEL J MANELLI
510(k) history	10 submissions · 10 cleared · 1992-2022

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k992404/> 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