

**K870518 SURGITEK PHYSICIAN DISPEN. PATIENT SELF-INJECT KIT**Apr 29, 1987  
83 days to decisionK870518 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k870518/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Feb 5, 1987
Decision date	Apr 29, 1987
Days to decision	83 days
Third-party review	No

**APPLICANT**

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Company	<b>Surgitek</b>
Location	Mchenry, IL, US
Contact	BETTY A LOCK
510(k) history	29 submissions · 28 cleared · 1983-1995

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k870518/>; 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 27, 2026