

**K900857 NON STERILE PARENTERAL SUPPLY KIT**May 22, 1990  
88 days to decisionK900857 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k900857/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - KD
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Feb 23, 1990
Decision date	May 22, 1990
Days to decision	88 days
Third-party review	No

**APPLICANT**

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Company	<b>Intermed, Inc.</b>
Location	Mchenry, IL, US
Contact	NORM RUEDT
510(k) history	16 submissions · 15 cleared · 1976-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900857/>; 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