

**K972705 RESHEATHER DEVICE-METAL OR PLASTIC,  
RESHEATER/ NEEDLE HOLDER DEVICE-PLASTIC**

Apr 30, 1998  
283 days to decision

K972705 · Product code: **FMI** · General Hospital  
Source: <https://www.510kdatabase.net/k972705/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jul 21, 1997
Decision date	Apr 30, 1998
Days to decision	283 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sage Products, Inc.</b>
Location	Mchenry, IL, US
Contact	KAREN PINTO
510(k) history	31 submissions · 30 cleared · 1978-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k972705/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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