

**K905783 MODIFIED WALKMED 400**Jan 11, 1991  
24 days to decisionK905783 · Product code: **FRN** · General Hospital  
Source: <https://www.510kdatabase.net/k905783/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Dec 18, 1990
Decision date	Jan 11, 1991
Days to decision	24 days
Third-party review	No

**APPLICANT**

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Company	<b>Medfusion, Inc.</b>
Location	Norcross, GA, US
Contact	RUFFIN BOOTH
510(k) history	5 submissions · 5 cleared · 1989-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k905783/>; 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 7, 2026