

**K893801 M.F.S. LOW AIR LOSS PULSATING BED**Jul 25, 1989  
64 days to decisionK893801 · Product code: **IOQ** · Physical MedicineSource: <https://www.510kdatabase.net/k893801/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bed, Flotation Therapy, Powered (IOQ)
Date received	May 22, 1989
Decision date	Jul 25, 1989
Days to decision	64 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Flotation Systems, Inc.</b>
Location	Herndon, VA, US
Contact	JACOBS, MD
510(k) history	5 submissions · 5 cleared · 1988-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k893801/>, 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 June 30, 2026