FDA 510(k) Premarket Notification

K893801 M.F.S. LOW AIR LOSS PULSATING BED

CLEARED

Jul 25, 1989
64 days to decision

K893801 · Product code: **IOQ** · Physical Medicine
Source: https://www.510kdatabase.net/k893801/

SUBMISSION DETAILS

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

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