

**K901295 PULMONAIR 40 PATIENT MANAGEMENT SYSTEM**Mar 30, 1990  
10 days to decisionK901295 · Product code: **IOQ** · Physical MedicineSource: <https://www.510kdatabase.net/k901295/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bed, Flotation Therapy, Powered (IOQ)
Date received	Mar 20, 1990
Decision date	Mar 30, 1990
Days to decision	10 days
Third-party review	No

**APPLICANT**

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Company	<b>The Medical Group, Inc.</b>
Location	Reno, NV, US
Contact	DIANE KOMOS
510(k) history	16 submissions · 16 cleared · 1986-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k901295/>; 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