FDA 510(k) Premarket Notification

K901295 PULMONAIR 40 PATIENT MANAGEMENT SYSTEM

CLEARED

Mar 30, 1990
10 days to decision

K901295 · Product code: **IOQ** · Physical Medicine
Source: https://www.510kdatabase.net/k901295/

SUBMISSION DETAILS

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

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