FDA 510(k) Premarket Notification

K882177 ADD-ON TO CRITICAL PATIENT CARE BED

CLEARED

Dec 1, 1988
191 days to decision

K882177 · Product code: **IOQ** · Physical Medicine
Source: https://www.510kdatabase.net/k882177/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bed, Flotation Therapy, Powered (IOQ)
Date received	May 24, 1988
Decision date	Dec 1, 1988
Days to decision	191 days
Third-party review	No

APPLICANT

Company	**Cardio Systems, Inc.**
Location	Mchenry, IL, US
Contact	CHARLES E HASTY
510(k) history	9 submissions · 9 cleared · 1979-2013