

**K882177 ADD-ON TO CRITICAL PATIENT CARE BED**Dec 1, 1988  
191 days to decisionK882177 · Product code: **IOQ** · Physical MedicineSource: <https://www.510kdatabase.net/k882177/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882177/>, 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 May 27, 2026