

**K860175 CD-200 CARDIAC MONITOR**Apr 15, 1986  
88 days to decisionK860175 · Product code: **DPS** · CardiovascularSource: <https://www.510kdatabase.net/k860175/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Jan 17, 1986
Decision date	Apr 15, 1986
Days to decision	88 days
Third-party review	No

**APPLICANT**

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Company	<b>Cardio Display Corp.</b>
Location	Westbury, NY, US
Contact	MILAN KESLER
510(k) history	2 submissions · 2 cleared · 1986-1986

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k860175/>; 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 July 7, 2026