

**K851325 MICROCOR ECG MONITOR**May 21, 1985  
47 days to decisionK851325 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k851325/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Apr 4, 1985
Decision date	May 21, 1985
Days to decision	47 days
Third-party review	No

**APPLICANT**

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Company	<b>Microcor, Inc.</b>
Location	Salt Lake City, UT, US
Contact	PERRY L LANE
510(k) history	2 submissions · 2 cleared · 1985-1986

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k851325/> 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