

**K872354 SERIES 2000 BEDSIDE PATIENT MONITOR**Jul 21, 1987  
33 days to decisionK872354 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k872354/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiometer & Rate Alarm) (DRT)
Date received	Jun 18, 1987
Decision date	Jul 21, 1987
Days to decision	33 days
Third-party review	No

**APPLICANT**

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Company	<b>Marquette Electronics, Inc.</b>
Location	Walker, MI, US
Contact	THOMAS MASSOPUST
510(k) history	82 submissions · 81 cleared · 1980-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k872354/>; 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 21, 2026