

**K895534 BIOLOG**Mar 19, 1990  
187 days to decisionK895534 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k895534/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Sep 13, 1989
Decision date	Mar 19, 1990
Days to decision	187 days
Third-party review	No

**APPLICANT**

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Company	<b>Micromedical</b>
Location	Gettysburg, PA, US
Contact	WESLEY KULP
510(k) history	2 submissions · 2 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895534/>; 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 6, 2026