

**K083534 LIFE BED NETWORK PATIENT VIGILANCE SYSTEM**Apr 13, 2009  
136 days to decisionK083534 · Product code: **DRT** · Cardiovascular  
Source: <https://www.510kdatabase.net/k083534/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiometer & Rate Alarm) (DRT)
Date received	Nov 28, 2008
Decision date	Apr 13, 2009
Days to decision	136 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hoana Medical, Inc.</b>
Location	Honolulu, HI, US
Contact	NANCY GERTLAR
510(k) history	5 submissions · 5 cleared · 2006-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k083534/> 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 June 21, 2026