

**K170973 VitalWatch Software User Interface**Jun 2, 2017  
60 days to decisionK170973 · Product code: **DRG** · CardiovascularSource: <https://www.510kdatabase.net/k170973/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Apr 3, 2017
Decision date	Jun 2, 2017
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vitalconnect, Inc.</b>
Location	Campbell, CA, US
Contact	Kevin Potgieter
510(k) history	10 submissions · 10 cleared · 2014-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k170973/> 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 27, 2026