

**K142672 AliveCor Heart Monitor**Dec 3, 2014  
75 days to decisionK142672 · Product code: **DXH** · Cardiovascular  
Source: <https://www.510kdatabase.net/k142672/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Transmitters And Receivers, Electrocardiograph, Telephone (DXH)
Date received	Sep 19, 2014
Decision date	Dec 3, 2014
Days to decision	75 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>AliveCor, Inc.</b>
Location	San Francisco, CA, US
Contact	Albert Boniske
510(k) history	19 submissions · 19 cleared · 2012-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k142672/> 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 24, 2026