

**K130921 ALIVECOR HEART MONITOR OTC**Feb 6, 2014  
309 days to decisionK130921 · Product code: **DPS** · Cardiovascular  
Source: <https://www.510kdatabase.net/k130921/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Apr 3, 2013
Decision date	Feb 6, 2014
Days to decision	309 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>AliveCor, Inc.</b>
Location	San Francisco, CA, US
Contact	MICHAEL RIGHTER
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/k130921/> 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