

**K173170 Epicardia Anywhere**Jan 10, 2018  
103 days to decisionK173170 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k173170/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Sep 29, 2017
Decision date	Jan 10, 2018
Days to decision	103 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medicomp, Inc.</b>
Location	Mchenry, IL, US
Contact	Sean Marcus
510(k) history	23 submissions · 23 cleared · 1983-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k173170/> 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 26, 2026