

**K171029 eCareCoordinator**Jul 14, 2017  
99 days to decisionK171029 · Product code: **DRG** · Cardiovascular  
Source: <https://www.510kdatabase.net/k171029/>**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 6, 2017
Decision date	Jul 14, 2017
Days to decision	99 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Visicu, Inc.</b>
Location	Lake Forest, CA, US
Contact	Daniel R. Plonski
510(k) history	7 submissions · 7 cleared · 2000-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k171029/> 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