

**K183322 physIQ Heart Rhythm and Respiratory Module**Jul 10, 2019  
222 days to decisionK183322 · Product code: **DPS** · Cardiovascular  
Source: <https://www.510kdatabase.net/k183322/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Nov 30, 2018
Decision date	Jul 10, 2019
Days to decision	222 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Physiq, Inc.</b>
Location	Naperville, IL, US
Contact	George Allen Hides
510(k) history	3 submissions · 3 cleared · 2018-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k183322/> 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 June 28, 2026