

**K172778 CheckMyHeart Plus**Jul 5, 2018  
294 days to decisionK172778 · Product code: **DPS** · CardiovascularSource: <https://www.510kdatabase.net/k172778/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Sep 14, 2017
Decision date	Jul 5, 2018
Days to decision	294 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Dailycare Biomedical, Inc.</b>
Location	Flagstaff, AZ, US
Contact	Ming Da Lee
510(k) history	5 submissions · 5 cleared · 2005-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k172778/> 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 30, 2026