

**K261286 The Circadia C300 System (C300)**May 20, 2026  
30 days to decisionK261286 · Product code: **DRT** · Cardiovascular  
Source: <https://www.510kdatabase.net/k261286/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Apr 20, 2026
Decision date	May 20, 2026
Days to decision	30 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Circadia Health, Inc.</b>
Location	El Segundo, CA, US
Contact	Erhan Ilhan
510(k) history	2 submissions · 2 cleared · 2026-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k261286/> 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 9, 2026