

**K172586 Vios Monitoring System™ Model 2050**Jun 22, 2018  
298 days to decisionK172586 · Product code: **DRT** · Cardiovascular  
Source: <https://www.510kdatabase.net/k172586/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Aug 28, 2017
Decision date	Jun 22, 2018
Days to decision	298 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vios Medical, Inc.</b>
Location	St. Paul, MN, US
Contact	Amit Patel
510(k) history	3 submissions · 3 cleared · 2015-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k172586/> 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