

**K240177 Zio AT® device (A100A1001)**Oct 30, 2024  
281 days to decisionK240177 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k240177/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jan 23, 2024
Decision date	Oct 30, 2024
Days to decision	281 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>iRhythm Technologies, Inc.</b>
Location	San Francisco, CA, US
Contact	Soyini Hamit
510(k) history	17 submissions · 17 cleared · 2008-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k240177/> 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 13, 2026