

**K233133 Talis EMR with Talis +ACG (Talis EMR with +ACG)**Apr 1, 2024  
187 days to decisionK233133 · Product code: **MWI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k233133/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Sep 27, 2023
Decision date	Apr 1, 2024
Days to decision	187 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Talis Clinical, LLC</b>
Location	Streetsboro, OH, US
Contact	William Murphy
510(k) history	1 submissions · 1 cleared · 2024-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k233133/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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