

**K142087 COVA MONITORING SYSTEM**May 1, 2015  
275 days to decisionK142087 · Product code: **MWI** · CardiovascularSource: <https://www.510kdatabase.net/k142087/>**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	Jul 30, 2014
Decision date	May 1, 2015
Days to decision	275 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Tosense, Inc.</b>
Location	San Diego, CA, US
Contact	NANDINI MIRTHY
510(k) history	2 submissions · 2 cleared · 2015-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k142087/> 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 1, 2026