

**K133810 VC150 VITAL SIGNS MONITOR**Oct 30, 2014  
318 days to decisionK133810 · Product code: **MWI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k133810/>**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	Dec 16, 2013
Decision date	Oct 30, 2014
Days to decision	318 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Innokas Yhtyma</b>
Location	Kempele, Oulu, FI
Contact	Tiina Kotipalo
510(k) history	1 submissions · 1 cleared · 2014-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k133810/> 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 5, 2026