

**K210204 Capsule Vitals Plus Patient Monitoring System**Oct 19, 2021  
266 days to decisionK210204 · Product code: **MWI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k210204/>**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	Jan 26, 2021
Decision date	Oct 19, 2021
Days to decision	266 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Capsule Technologie Sas</b>
Location	Andover, MA, US
Contact	Peter Kelley
510(k) history	4 submissions · 4 cleared · 2015-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k210204/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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