

**K192137 IntelliVue Patient Monitor MX500, IntelliVue Patient Monitor MX550**

Sep 11, 2020  
401 days to decision

K192137 · Product code: **MHX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k192137/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Aug 7, 2019
Decision date	Sep 11, 2020
Days to decision	401 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Philips Medizin Systeme Boeblingen GmbH</b>
Location	B?blingen, DE
Contact	Stefan Breuer
510(k) history	48 submissions · 48 cleared · 2004-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k192137/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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