

**K251702 IntelliVue Patient Monitor MX750**

Nov 10, 2025  
161 days to decision

K251702 · Product code: **MHX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k251702/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Jun 2, 2025
Decision date	Nov 10, 2025
Days to decision	161 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	IntelliVue Patient Monitor MX850; IntelliVue 4-Slot Module Rack FMX-4 (866471 866470 866468 )

**APPLICANT**

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Company	<b>Philips Medizin Systeme B?blingen GmbH</b>
Location	Boeblingen, DE
Contact	Alicia Honemeyer
510(k) history	8 submissions · 8 cleared · 2022-2025

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

Device record: <https://www.510kdatabase.net/k251702/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).

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