

K221348 IntelliVue Patient Monitor MX750 (866471)Feb 3, 2023
270 days to decisionK221348 · Product code: **MHX** · CardiovascularSource: <https://www.510kdatabase.net/k221348/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	May 9, 2022
Decision date	Feb 3, 2023
Days to decision	270 days
Third-party review	No
Summary / Statement	Summary
Other names	IntelliVue Patient Monitor MX850 (866470)

APPLICANT

Company	Philips Medizin Systeme B?blingen GmbH
Location	Boeblingen, DE
Contact	Stefan Breuer
510(k) history	8 submissions · 8 cleared · 2022-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k221348/> Data sourced from FDA 510(k) public records (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. - Generated May 24, 2026