

K181314 IntelliVue Patient Monitor MX100, IntelliVue Multi-Measurement Module X3, IntelliVue Microstream ExtensionSep 4, 2018
110 days to decisionK181314 · Product code: **MHX** · Cardiovascular
Source: <https://www.510kdatabase.net/k181314/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	May 17, 2018
Decision date	Sep 4, 2018
Days to decision	110 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k181314/>; 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 18, 2026