FDA 510(k) Premarket Notification

CLEARED

K221348 IntelliVue Patient Monitor MX750 (866471)

Feb 3, 2023
270 days to decision

K221348 · Product code: **MHX** · Cardiovascular
Source: 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