FDA 510(k) Premarket Notification

CLEARED

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

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

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