

**K061052 THE PHILIPS INTELLIVUE PATIENT MONITORS,  
MODELS MP20,MP30,MP40,MP50,MP60,MP70,MP80,MP90**

May 26, 2006  
39 days to decision

K061052 · Product code: **MHX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k061052/>

**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Apr 17, 2006
Decision date	May 26, 2006
Days to decision	39 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Philips Medical Systems North America, Inc.</b>
Location	Shelton, CT, US
Contact	HAUKE SCHIK
510(k) history	71 submissions · 71 cleared · 1989-2010

---

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

Device record: <https://www.510kdatabase.net/k061052/> 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 17, 2026