

**K032858 THE PHILIPS INTELLIVUE MP40, MP50, MP60,
MP70, AND MP90 PATIENT MONITORS, RELEASE B.O.**Oct 10, 2003
28 days to decisionK032858 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k032858/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Sep 12, 2003
Decision date	Oct 10, 2003
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Philips Medizin Systeme Boblingen GmbH
Location	Andover, MA, US
Contact	HAUKE SCHIK
510(k) history	12 submissions · 12 cleared · 2001-2015

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k032858/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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