

**K131913 INTELLIVUE CL SPO2 POD AND CL NBP POD,
PHILIPS INTELLIVUE PATIENT MONITORS MP5, MP5T, MP5SC**Nov 26, 2013
153 days to decisionK131913 · Product code: **DRG** · Cardiovascular
Source: <https://www.510kdatabase.net/k131913/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Jun 26, 2013
Decision date	Nov 26, 2013
Days to decision	153 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k131913/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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