

**K122223 NEW: INTELLIVUE CL RESPIRATION POD NEW:
SPECIALIZED ASSESSORY: MOBILE CL RESP ATTACHMENT
MODIFIED: INTELLIVUE PATIENT MO**Apr 12, 2013
261 days to decisionK122223 · Product code: **BZQ** · Anesthesiology
Source: <https://www.510kdatabase.net/k122223/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Breathing Frequency (BZQ)
Date received	Jul 25, 2012
Decision date	Apr 12, 2013
Days to decision	261 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/k122223/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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