FDA 510(k) Premarket Notification

K131829 INTELLIVUE PATIENT MONITOR

CLEARED

Aug 16, 2013
57 days to decision

K131829 · Product code: **MHX** · Cardiovascular
Source: https://www.510kdatabase.net/k131829/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Jun 20, 2013
Decision date	Aug 16, 2013
Days to decision	57 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Philips Medizinsysteme Boeblingen GmbH, Cardiac AN**
Location	Boeblingen, Baden-Wuerttemberg, DE
Contact	HERBERT VAN DYK
510(k) history	25 submissions · 25 cleared · 2007-2014

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k131829/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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