

K082280 SURESIGNS VS2 VITAL SIGNS MONITOR, SURESIGNS VM1 PATIENT MONITOR, SURESIGNS VS3 CITAL SIGNS MONITOROct 8, 2008
58 days to decisionK082280 · Product code: **MHX** · Cardiovascular
Source: <https://www.510kdatabase.net/k082280/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Aug 11, 2008
Decision date	Oct 8, 2008
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Philips Medical Systems North America Co.
Location	Shelton, CT, US
Contact	MARY KRUITWAGEN
510(k) history	24 submissions · 24 cleared · 2001-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k082280/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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