FDA 510(k) Premarket Notification

K123900 SURESIGNS VM4, SURESIGNS VM6, SURESIGNS VM8

CLEARED

Apr 18, 2013
121 days to decision

K123900 · Product code: **MHX** · Cardiovascular
Source: https://www.510kdatabase.net/k123900/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Dec 18, 2012
Decision date	Apr 18, 2013
Days to decision	121 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Philips Medical Systems**
Location	Seattle, WA, US
Contact	MARY KRUITWAGEN
510(k) history	107 submissions · 105 cleared · 2002-2021

Philips Medical Systems is a Dutch multinational health technology company headquartered in Amsterdam with U.S. operations based in Seattle. The company evolved from a consumer electronics conglomerate founded in 1891 to a healthcare-focused organization. Philips Medical Systems has received FDA 510(k) clearances from total submissions between 2002 and 2021. The company's regulatory focus centered on Cardiovascular devices, which represented 79% of all submissions. This historical record reflects the company's significant presence in diagnostic ultrasound systems and pati...

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k123900/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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