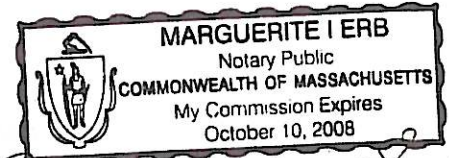


CE**DECLARATION OF CONFORMITY****Manufacturer's Name:** Philips Medical Systems**Manufacturer's Address:** 3000 Minuteman Road
Andover, MA 01810-1099
United States**Declares, that the product****Product Name:** SureSigns Hardware Bundle**Model Number(s):** NA**Product Options/Accessories:** *Standard Accessories*
router
hub
mouse*One of the following keyboards*
US English or equivalent
Spanish or equivalent
French or equivalent
German or equivalent
Dutch or equivalent
Norwegian or equivalent
Italian or equivalent
Danish or equivalent
Swedish or equivalent*Optional accessories*
Li-ion Battery pack
Wall mount

Marguerite I ERB

**to which this declaration relates is in conformity with the European Council Directive:
EMC Directive 2004/108/EEC
and carries the CE-marking accordingly.**


Supplementary Information:

The individual devices comply with the requirements of the EMC Directive 204/108/EEC and where appropriate comply with the requirements of the Low Voltage Directive 2006/95/EC and carry CE-marking by their respective manufacturers. The product (system) was tested in a typical configuration as described in the Manufacturer's accompanying documents. The testing configuration includes the SureSigns VSV, the hub, the mouse and a keyboard. They also include the recorder and paper, the battery and the second display.

Andover, 27 September 2007

Denise Haley
Denise Haley,
Senior Manager, Regulatory Affairs
Patient Monitoring
Cardiac and Monitoring Systems
Philips Medical Systems

Authorized EU-representative: Philips Medizin Systeme Böblingen GmbH, Hewlett-Packard Str. 2, 71034 Böblingen, Germany

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