OPIS

CE 0

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: IntelliVue 2.4 GHz Smart Hopping Access Point
(A component of the ITS4850A 2.4 GHz IntelliVue Telemetry System)

Model Number(s): ITS4852A

Product Options/Accessories: None.

to which this declaration relates is in conformity with the European Council Directives:
Low Voltage Directive 2006/95/EC,
EMC Directive 89/336/EEC
And
R&TTE Directive 1999/5/EC
and carries the CE-marking accordingly.

Supplementary Information:
The product was tested in a typical configuration as described in the Manufacturer's accompanying documents. The product is a component of the ITS4850A 2.4 GHz IntelliVue Telemetry System.

The radio transmitting equipment used in this product is Class 2 under the scope of the R&TTE Directive and was tested according to Article 10 via Annex III, and all essential radio test suites (as defined in the Essential Requirements) have been carried out.
This product is intended to be connected to the Publicly Available Interfaces (PAI) and used throughout the EEA. Individual countries may apply restrictions on putting this device into service or placing on the market.

MARGUERITE I ERB
Notary Public
COMMONWEALTH OF MASSACHUSETTS
My Commission Expires
October 10, 2008

A True Copy Attest
MARGUERITE I. ERB
NOTARY PUBLIC
My commission expires Oct. 10, 2008

Andover, 26 February 2007

L.S.

Peter Ohanian, Director, Quality and Regulatory Affairs
Patient Monitoring - Ultrasound and Monitoring
Philips Medical Systems

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