ZERTIFIKAT ◆ CERTIFICATE ◆ 证书 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Product Service

EC-CERTIFICATE

Full Quality Assurance System

(Annex II, section 3 of the Directive 93/42/EEC on Medical Devices)

No. G1 09 04 44649 007

Manufacturer:

Philips Medical Systems
3000 Minuteman Road
Andover MA 01810-1099
USA

EC-Representative:

Philips Medizin Systeme Böblingen GmbH
Hewlett-Packard Strasse 2
71043 Böblingen
GERMANY

Product Category(ies):

Single and Multi-Parameter Patient Monitors (Transport, Point-of-Care, Central Unit, Home Use); Cardiographs; ICG Systems; ECG Systems; EEG Systems; Telemetry Systems; Defibrillators and related Accessories (including Cables, Electrodes, Recorders) for ECG, Cardiac Output, ICG, EEG, CPR Feedback, Oxygen Saturation, Respiration and Gas Monitoring, Weight, Blood Pressure (invasive and Non-Invasive), Temperature, Physiological Pressure Kits, Defibrillation (including Paddles) and Clinical Information Systems; Transesophageal Sheath Kits

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective products / product categories according to Annex II section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system conforms to the provisions of the Directive and is subject to periodical surveillance. For marketing of class II products an additional Annex II 4 certificate is mandatory. See also notes overleaf.

Report No DM90017

Valid until 2014-08-05

Date, 2009-08-06

Signed / Stamped / Authorised

TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123