ZERTIFIKAT ◆ CERTIFICATE ◆ 認証証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Product Service

EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 16 02 78838 006

Manufacturer:

Philips Medical Systems
22100 Bothell Everett Highway
Bothell WA 98021
USA

EC-Representative:

Philips Medizin Systeme Böblingen GmbH
Hewlett-Packard Strasse 2
71034 Böblingen
GERMANY

Product Category(ies):

External Defibrillator Systems and Defibrillation Electrodes

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 devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: 72109937

Valid from: 2016-07-06
Valid until: 2021-07-05

Date, 2016-05-27

Stefan Preiß

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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Facility(ies):

Philips Medical Systems
22100 Bothell Everett Highway, Bothell WA 98021, USA

Philips Medical Systems
21919 20th Avenue SE, Bothell WA 98021-8431, USA

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