ZERTIFIKAT ◆ CERTIFICATE ◆ 認証書 ◆ 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 15 06 19931 008

**Manufacturer:**

**GS Elektromedizinische Geräte G. Stemple GmbH**
Hauswiesenstr. 26
86916 Kaufering
GERMANY

**Facility(ies):**

GS Elektromedizinische Geräte G. Stemple GmbH
Hauswiesenstr. 26, 86916 Kaufering, GERMANY

**Product Category(ies):**

**Defibrillators, patient monitoring systems, resuscitation equipment, Software for the Transmission of medical data**

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.:** 713064068

**Valid from:** 2015-11-08
**Valid until:** 2020-11-07

**Date,** 2015-11-06

Hans-Heiner Junker


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





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