



Benannt durch/Designated by
 Zentralstelle der Länder
 für Gesundheitsschutz
 bei Arzneimitteln und
 Medizinprodukten
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 ZLG-BS-244.10.08



Product Service

EC Certificate

Production Quality Assurance System
 Directive 93/42/EEC on Medical Devices (MDD), Annex V
 (Devices in Class IIa, IIb or III)

No. G2 011882 0080 Rev. 00

Manufacturer: **PHYSIO-CONTROL, Inc.**
 11811 Willows Road N.E.
 Redmond WA 98052
 USA

Product Category(ies): **Cardiac Defibrillator Accessories**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

Report No.: 72151772

Valid from: 2020-03-17
Valid until: 2024-05-26

Date, 2020-03-17

Christoph Dicks
 Head of Certification/Notified Body

TÜV SÜD
 ZERTIFIKAT ◆ CERTIFICATE ◆ 認證書 ◆ CERTIFICADO ◆ CERTIFIKAT ◆ CERTIFICATE