



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



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 079546 0023 Rev. 00

Manufacturer:

ZOLL Medical Corporation

269 Mill Road
Chelmsford MA 01824-4105
USA

EC-Representative:

Zoll International Holding B.V.
Newtonweg 18, 6662 PV Elst, THE NETHERLANDS

Product Category(ies):

**External Defibrillators, Automatic External
Defibrillators, Multi-Parameters Monitoring
Systems, Defibrillation Electrodes, Blood
Cartridge and Crystalloid/Colloid Cartridge,
Portable Ventilators with Patient Breathing
Circuits.**

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.:

72145841

Valid from:

2019-07-30

Valid until:

2024-05-26

Date,

2019-07-30

Stefan Preiß
Head of Certification/Notified Body



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Facility(ies):

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