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

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

No. G2 055329 0025 Rev. 00

Manufacturer:

Product Category(ies):

Electric Suction Apparatus,
Oxygen Concentrator,
Air Compressive Nebulizer,
Electronic Blood Pressure Monitor,
Finger Pulse Oximeter, Non-contact Infrared Forehead Thermometer,
Portable Phlegm Suction Unit
Mesh Nebulizer

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.: SH2039201

Valid from: 2020-03-30
Valid until: 2024-05-26

Date, 2020-03-30

Christoph Dicks
Head of Certification/Notified Body

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

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