



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 003389 0002 Rev. 01

Manufacturer: **Silbermann Technologies Ltd.**
 5 HaRakevet Street
 4900715 Petach Tikva
 ISRAEL

Product Category(ies): **Medical Gas Supply Systems and Oxygen
 and Vacuum Therapy Units,
 Components and Accessories**

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.: 713151907

Valid from: 2020-03-02
Valid until: 2023-07-09

Date, 2020-03-02

Christoph Dicks
 Head of Certification/Notified Body