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 051287 0013 Rev. 01

Manufacturer: **Herrmann Apparatebau GmbH**
Im Höning 3
63820 Elsenfeld
GERMANY

Facility(ies): Herrmann Apparatebau GmbH
Im Höning 3, 63820 Elsenfeld, GERMANY

Product Category(ies): **Colon-Therapy Devices including Disposables and Ozone-Therapy-Devices**

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.: 713139922

Valid from: 2018-10-18
Valid until: 2023-10-17

Date, 2018-10-11

Stefan Preiß

Digitally signed by Şarco Alexandru
Date: 2023.03.10 19:56:36 EET
Reason: MoldSign Signature
Location: Moldova

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

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A4 / 07.17