EU Quality Management System Certificate

mdc medical device certification GmbH

Kriegerstr. 6, 70191 Stuttgart, Germany
Notified body (identification number 0483)

hereby certifies that the company (SRN: LI-MF-000002555)

BEMER Int. AG

Austrasse 15
9495 Triesen
Liechtenstein

has implemented and applies a quality management system in accordance with Annex IX, Chapter I of Regulation (EU) 2017/745 for conformity assessment of the devices listed on the following pages.

An audit by mdc has proven that this quality management system fulfils the following requirements:

Annex IX - Chapter I (Quality Management System)

of Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices

Surveillance is carried out in accordance with Annex IX, Section 3 of Regulation (EU) 2017/745.

This certificate consists of 2 pages. Details of the devices affected by this certificate as well as further information and conditions are included on the following pages.

Valid from:	2023-02-15	Registration No.	D1304700012
Valid until:	2027-10-28	Evaluation Report No.	P22-00069-225696

Stuttgart, 2023-02-15

Head of Notified Body

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
BS-MDR-098

Devices:

Product:
stimulation devices for physical vascular therapy

- Control unit [B.Box Evo]
- Local Applicator [B.Spot Evo; B.Pad Evo; B.Sit Evo]
- Full Body Applicator [B.Body Evo; B.Bed Evo]
- Light Applicator [B.Light Clear Evo; B.Light Restore Evo]

Risk class: IIa