EU Quality Management System Certificate

We hereby certify the company

Hans Dinslage GmbH
Riedlinger Straße 28
88524 Uttenweiler
Germany

the introduction and application of a quality management system in accordance with Annex IX, Chapter I and III of Regulation (EU) 2017/745 for conformity assessment.

An audit by mdc has proven that this quality management system meets 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 from mdc medical device certification GmbH (Notified Body 0483) consists of 3 pages. Details about the devices covered as well as further information and conditions are contained on the following pages.

Valid from	2026-04-09	Registration No.	D1326900052
Valid until	2030-11-21	Report No.	P25-00469-359089

Stuttgart, 2026-04-08

Notified Body

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

Devices:

Pulse oximeters

Risk class: IIa

Blood pressure monitors

Risk class: IIa

Thermometers

Risk class: IIa

Bite healers

Risk class: IIa

Massage devices

Risk class: IIa

Hearing aids

Risk class: IIa

Infrared lamps

Risk class: IIa

Nebulizers

Risk class: IIa

TENS devices

Risk class: IIa

The certificate is based on the previous certificate

D1326900036 (2021-04-09)
D1326900039 (2021-11-22)
D1326900041 (2022-05-20)
D1326900042 (2023-03-14)
D1326900045 (2023-12-14)
D1326900048 (2025-02-07)
D1326900049 (2025-10-13)

with the following changes to D1326900049:
Re-Certification with new term