People.Health.Care.

EU Declaration of Conformity DOC-0002-04

We declare under our sole responsibility that the following products comply with Regulation (EU) 2017/745 **(MDR)** on medical devices.

Manufacturer	**Lohmann & Rauscher International GmbH & Co. KG** Westerwaldstraße 4 56579 Rengsdorf Germany
SRN	DE-MF-000005052
Product Group	**Short Stretch Compression Bandages**
Catalogue numbers & product names	See document DI_REF_00021_02
Intended Purpose	For compression, retention of dressings, support and relief and for immobilisation of limbs.
Basic UDI-DI	4021447-0002-JQ (sterile) 4021447-0002A-NST-XL, 4021447-0002B-NST-XZ, 4021447-0002C-NST-YE, 4021447-0002D-NST-YT (non-sterile, see DI_REF_00021_02)
Class / Rule	I/01 + Is/01
Conformity Assessment Route	Class 1 non-sterile variants: Technical documentation according to annexes II and III of Regulation (EU) 2017/745 Class 1 sterile variants: Annex XI Part A of Regulation (EU) 2017/745
Notified Body:	Only applicable for Basic UDI-DI 4021447-0002JQ

TÜV SÜD Product Service GmbH
Ridlerstraße 65
80339 Munich
Germany
Identification number: 0123

Certificate identification:
G21 045286 0081

Device Group on Certificate:
M03 BANDAGES

Oliver Opitz

Oliver Opitz, Mar 27, 2026 09:04 UTC

Neuwied

Place

Lohmann & Rauscher International GmbH & Co. KG
Oliver Opitz, PRRC-Q