People.Health.Care.

EU DECLARATION OF CONFORMITY DOC-0030-05

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	**Tubular bandages**
Catalogue numbers & product names	See document DI_REF_00103_02
Intended Purpose	Tubular bandages are intended for retention of wound dressings and as skin protection under Plaster of Paris (P.o.P.), synthetic casts and compression bandages.
Basic UDI-DI	4021447-0030-JZ (sterile); 4021447-0030NST-4M (non-sterile)
Class / Rule	I/01 + Is/01
Conformity Assessment Route	non sterile variants: Technical documentation according to annexes II and III of Regulation (EU) 2017/745 sterile variants: Annex XI Part A of Regulation (EU) 2017/745
Notified Body:	Only applicable for Basic UDI-DI 4021447-0030-JZ

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

Certificate identification:
G21 045286 0081 Annex XI Part A

Device Group on Certificate:
M03 - BANDAGES

Oliver Opitz

Oliver Opitz, Jun 09, 2026 07:51 UTC

Neuwied
Place

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