

**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, Jun 09, 2026 07:51 UTC

Neuwied

Place

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