



EU Declaration of Conformity

Heidenheim, 2025-03-25

Product Group Number	2122
Manufacturer	PAUL HARTMANN AG Paul-Hartmann-Str. 12 89522 Heidenheim GERMANY

We herewith declare under our sole responsibility that the product listed above, first placed on the market by PAUL HARTMANN AG, satisfies the applicable provisions of the following listed legislation. The conformity assessment procedure has been performed and the Technical Documentation is kept available.

Applied legislative act

Medical Device Regulation (EU) 2017/745

High Level Intended Purpose	Non-active, non-implantable devices for wound and skin care	
Classification	Risk Class	Rule
	Class I	Rule 1
Conformity Assessment Procedure	Article 52 (7)	
Notified Body	n/a	
Single Registration Number	Manufacturer: DE-MF-000005861	
Basic UDI-DI	40495002122JV	
EMDN	Code	Term
	M030401	ELASTIC COMPRESSION BANDAGES
GMDN	61223	Pressure bandage, non-latex, non-sterile, reusable
UMDNS	0	0

GLN 404 9500 00000 0

Vorstand/Management Board: Britta Fünfstück
 (Vorsitzende des Vorstands/CEO), François Georgelin,
 Stefan Grote, Oliver Neubrand
 Aufsichtsratsvorsitzender/Chairman of the Supervisory Board:
 Fritz-Jürgen Heckmann

Sitz Heidenheim
 Amtsgericht Ulm HRB 661090
 Registered Office Heidenheim
 Commercial Register of the District Court of Ulm file no. HRB
 661090

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Helps. Cares. Protects.

PAUL HARTMANN AG

ppa.

Jochen Bauer

Vice President R&D Wound Care & Consumer
Health

ppa.

Stefan Fischer

Senior Vice President Regulatory Affairs

Valid until: 2030-10-31

List of products falling under the respective Product Group Number:

REF	Name - Description
931825	Puetter flex 8/10cm x 5m P2
931826	Puetter flex duo 10cmx5m P2
931827	Puetter flex 6cmx5m P1
931828	Puetter flex 8cmx5m P1
931829	Puetter flex 10cmx5m P1
931830	Puetter flex 12cmx5m P1
931832	Puetter flex 8cmx5m P10
931833	Puetter flex 10cmx5m P10

GLN 404 9500 00000 0

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