



EU Declaration of Conformity

Heidenheim, 2024-10-31

Product Group Number	1278
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	TÜV SÜD Product Service GmbH, Ridlerstr. 65, 80339 München, Germany, ID-Nr. 0123	
Single Registration Number	Manufacturer: DE-MF-000005861	
Basic UDI-DI	40495001278KN	
EMDN	Code	Term
	M030401	ELASTIC COMPRESSION BANDAGES
GMDN	61223	Pressure bandage, non-latex, non-sterile, reusable
UMDNS	15-557	Binde (Binders)

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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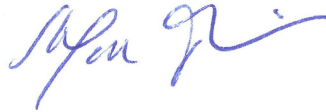


PAUL HARTMANN AG

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Jochen Bauer
Vice President R&D Wound Care & Consumer
Health



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
935151	Hospilite 5cm x 4,5m cel
935152	Hospilite 7,5cm x 4,5m cel
935153	Hospilite 10cm x 4,5m cel
935154	Hospilite 15cm x 4,5m cel

GLN 404 9500 00000 0

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