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## EU Declaration of Conformity Class I

Heidenheim, 2024-09-24

We herewith declare under our sole responsibility that the Class I medical devices listed below, first placed on the market by PAUL HARTMANN AG, Single Registration Number of Manufacturer DE-MF-000005861, satisfy the applicable provisions, in particular, the General Safety and Performance Requirements, of the Regulation (EU) 2017/745 of the European Parliament and of the Council of 5. April 2017 on medical devices.

The conformity assessment procedures according to Article 52 (7) have been performed and the Technical Documentation is kept available.

Intended Purpose	Non-active, non-implantable devices for incontinence care, worn on the body		
Product Name	Product Group Number	Classification Rule (according to Annex VIII of Regulation (EU) 2017/745)	Basic UDI-DI
MoliCare Premium Bed Mat	3212	1	40495003212K6
Hartmann Bed Mat	3212	1	40495003212K6

PAUL HARTMANN AG

**Stefan Grote**  
Member of the Management Board

ppa.

**Stefan Fischer**  
Senior Vice President Regulatory Affairs

Valid until: 2025-06-30

GLN 404 9500 00000 0

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(Vorsitzende des Vorstands/CEO), François Georgelin,  
Stefan Grote, Stefan Müller  
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Sitz Heidenheim  
Amtsgericht Ulm HRB 661090  
Registered Office Heidenheim  
Commercial Register of the District Court of Ulm file no. HRB  
661090