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HARTMANN
Helps. Cares. Protects.

UK Declaration of Conformity for Medical Devices in Class I unsterile

Heidenheim, 2025-05-28

We herewith declare under our sole responsibility of the manufacturer that the Class I medical device 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 Essential Requirements, of UK Medical Devices Regulations 2002 (SI 2002 No 618, as amended) which were amended by the Medical Devices (EU Exit) Regulations 2019 and the Medical Devices (EU Exit) Regulations 2020.

The conformity assessment procedures according to Annex VII have been performed and the Technical Documentation is kept available.

UK Representative

PAUL HARTMANN LTD,

Unit P2, Parklands, Heywood Distribution Park, Pilsworth Road

Heywood, Lancs., OL10 2TT

Intended Purpose	Non-active, non-implantable devices for incontinence care, worn on the body		
Product Name	**Product Group Number**	**Classification Rule** (according to Annex IX of Directive 93/42)	**Basic UDI-DI**
MoliCare Premium Men Pad	3122	1	40495003122K4

PAUL HARTMANN AG

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Stefan Grote
Member of the Management Board

Stefan Fischer
Senior Vice President Regulatory Affairs

Valid until: 2026-05-28

GLN 404 9500 00000 0
Vorstand/Management Board: Britta Fünfstück (Vorsitzende des Vorstands/CEO), François Georgelin, Stefan Grote, Stefan Müller
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