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HARTMANN
Helps. Cares. Protects.

EU Declaration of Conformity Class I

Heidenheim, 2024-10-07

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 Mobile XXL (ADL)	4009	1	40495004009KE

PAUL HARTMANN AG

i.V.

Stefan Grote
Member of the Management Board

ppa.

Stefan Fischer
Senior Vice President Regulatory Affairs

Valid until: 2025-06-30