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## EU Declaration of Conformity Class I

Heidenheim, 2024-02-13

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 Pad     | 3120  | 1   | 40495003120JY |
| HARTMANN Pad     | 3120  | 1   | 40495003120JY |

PAUL HARTMANN AG

**Stefan Grote**  
Member of the Management Board

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**Stefan Fischer**  
Senior Vice President Regulatory Affairs

Valid until: 2024-06-30

GLN 404 9500 00000 0

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(Vorsitzende des Vorstands/CEO), François Georgelin,  
Stefan Grote, Oliver Neubrand

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