Medical Device Declaration of Conformity

Version:2	Revision Date: 13.06.2025	Date of First Issue: 10.06.2022	TD:00125

Supplier and Product Details	
Manufacturer / Place of Issue	Kimberly-Clark B.V. Copernicuslaan 35, 6716 BM Ede, Netherlands
Product Name	DryNites Bed Mats
Product Codes	2907049, 2907053, 2043905
Competent Authority	CIBG, Ministry of Health, Welfare and Sport, Netherlands
SRN	NL-MF-000004367
EMDN	T04010201 Untackable Underpads
Classification Rule	One
Basic UDI-DI	8710461BEDMATSPS

This EU declaration of conformity is issued under the sole responsibility of Kimberly Clark B.V. being a manufacturer.

We hereby declare that the above mentioned devices comply with the European Medical Devices Regulation 2017/745, and are in accordance with Article 52-7 of Medical Devices Regulation 2017/745, supported by the Conformity Assessment Procedure, and adhering to the general safety and performance requirements in accordance with Annex I of Regulation 2017/745. This declaration is made on the basis of an implemented documented Quality Management System, and is also based on the existing Technical Documentation as per Annex II and III. CE marking carried out as per Annex V of Regulation 2017/745. This declaration is valid for the above product in its original, unmodified packaging.

Ozlem Ariz Sr Regulatory Affairs Manager, Europe		Revision Date: 13.06.2025

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