

EUROPEAN MEDICAL DEVICE REGULATION**Declaration of Conformity**

As Legal Manufacturer, we

Solventum Germany GmbH

Edisonstrasse 6,

59174 Kamen, Germany

Single Registration Number: DE-MF-000038373

hereby declare under our sole responsibility that the following CE marked devices

Trade Name	Solventum™ Medipore™ +Pad Adhesive Wound Dressing
Intended Purpose	Adhesive wound dressing is designed to be used as a mechanical barrier and for absorption of exudates for acute wounds.
Reference	3562E, 3564E, 3566E, 3569E, 3570E, 3571E, 3573E 3562NP, 3566NP, 3569NP, 3570NP, 3562SP, 3566SP, 3562P-10, 3566P-10, 3569P-10, 3570P-10, 3562PR-10, 3566PR-10
Basic UDI-DI	01979982761010000000032PZ

are classified per rule 4 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class 1 sterile devices in accordance with Annex IX and all other applicable provisions of the REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL concerning medical devices.

This declaration is made based on the quality assurance certificate
EU Certificate Number: 31619899 MDR2017Q
Issued by: DQS Medizinprodukte GmbH, No. 0297

DocuSigned by:

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30-Jun-2025 | 16:56 CEST

Claudia Inden
Senior Manager Regulatory Affairs
Medical Surgical Business (MedSurg)
Solventum Germany GmbH

Date