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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 device(s)

Trade Name	**Solventum™ Tegaderm™ + Pad Film Dressing with Non-Adherent Pad**
Intended Purpose	Dressing is designed for covering superficial and partial-thickness wounds, surgical incisions and IV catheter sites.
Reference	3582E, 3584E, 3586E, 3589E, 3590E, 3591E, 3593E
Basic UDI-DI	01979982761010000000253QK

are classified per rule 4 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class IIa 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 certificate
EU Quality Management Certificate: 31619899 MDR2017Q
Issued by: DQS Medizinprodukte GmbH, August-Schanz-Straße 21, 60433 Frankfurt am Main, Germany No. 0297

DocuSigned by:
Claudia Inden
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22-Jul-2025 | 08:42 CEST

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

Date