EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

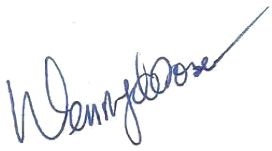
3M Deutschland GmbH
Single Registration Number DE-MF-000011641
Carl-Schurz-Str. 1
41453 Neuss
Germany

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

Trade Name	Nexcare™ Blood Stop
Intended Purpose	Nexcare™ Blood Stop is a non-invasive device intended to come into contact with injured skin
Reference	N1714AS, N1730AS, N1714NS, BS-14
Basic UDI-DI	06082232761050000000022GH

is classified per rule 4 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class IIa 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
EC Certificate Number: 003626 MDR2017Q
Issued by: DQS Medizinprodukte GmbH, No. 0297



22 December 2025

Wenny Sumarni
EMEA Regulatory Affairs Manager
3M Consumer Health Care

Date

Related to REG-STED-MDR-05-731306