EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we
3M Deutschland GmbH
Health Care Business
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	3M™ Kerracel™ Gelling Fiber Dressing
Intended Purpose	3M™ Kerracel™ Gelling Fiber Dressing is designed for the management of moderately to heavily exuding partial and full-thickness chronic wounds and acute wounds, and to control minor bleeding in superficial wounds.
Reference	CWL1032, CWL1033, CWL1034, CWL1035, CWL1166, CWL1032F, CWL1166F, CWL1237F, CWL1238F
Basic UDI-DI	06082238401010000000201A6

is classified per rule 4 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class IIb device 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 EU Quality Management Certificate:

EU Quality Management Certificate: 003626MDR2017Q

Issued by: DQS Medizinprodukte GmbH, August-Schanz-Straße 21, 60433 Frankfurt am Main, Germany, No. 0297

Harald Ceschinski
Manager Regulatory Affairs and
Quality Management System
Health Care Business EMEA
3M Deutschland GmbH

Neuss, February 2, 2024
Location/Date