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# *Declaration of Conformity*

We
**3M Deutschland GmbH**
**Health Care Business**
**Carl-Schurz-Str. 1**
**41453 Neuss**
**Germany**

hereby declare under our sole responsibility
that the CE marked products, to which this declaration relates,

**Product Name**
**3M™ Kerramax Care™ Super-Absorbent Dressing**

**Ref Code**
**PRD500-050, PRD500-120, PRD500-240, PRD500-380-B10, PRD500-025, PRD500-600-B10, PRD500-100, PRD500-050-B50, PRD500-120-B50, PRD500-240-B30, PRD500-380-B30, PRD500-025-B550, PRD500-050-B550, PRD500-100-B550, PRD500-065**

are classified per rule 4 of Annex IX of the Medical Device Directive 93/42/EEC,
as **Class IIb** devices
and

are in accordance with
***Annex II and all other applicable provisions of the Directive 93/42/EEC***
on the approximation of the laws of the European Union Member States concerning medical devices.

This declaration is made on the basis of the quality assurance certificate
EC Certificate No. 003626 MR2 valid until 2024-05-26 delivered by DQS Medizinprodukte GmbH,
August-Schanz-Straße 21, D-60433 Frankfurt am Main, No. 0297

Signature Margaret Bessenbach July 12, 2021

Margaret Bessenbach
Manager Regulatory and Quality
Health Care Business EMEA
3M Deutschland GmbH

Neuss / Date