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EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Company
Single Registration Number: US-MF-000014086
2510 Conway Ave. St. Paul, MN 55144 USA

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

Trade Name	Nexcare™ Liquid Bandage Spray MPlast Sprühpflaster
Intended Purpose	A unique skin shield (protectant) that doesn't sting when applied. It is alcohol-free and soothes minor skin scrapes and abrasions. When applied, the thin shield provides breathable, waterproof protection that covers and protects skin to allow healing.
Reference	LBS-118 5203.528
Basic UDI-DI	06082238401050000000047EG

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

EU Authorized Representative:

3M Deutschland GmbH
Health Care Business
DE-AR-000011642
Carl-Schurz-Str. 1
41453 Neuss, Germany

DocuSigned by:
Bryan Becker
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9/29/2022

Bryan Becker
Director, Quality Assurance and Regulatory Affairs
3M Consumer Health and Safety Division
2510 Conway Ave. St. Paul, MN 55144 USA

Date

3M and Nexcare are trademarks of 3M.