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™ Steri-Strip™ First Aid Skin Closures Viscoplast™ Steri-Strip™ First Aid Skin Closures Spofaplast® Steri-Strip™ First Aid Skin Closures M-Plast Wundverschluss-Streifen
Intended Purpose	Intended to secure, close, and support small cuts and wounds. To be used as directed by physician or health care provider for treatment of small cuts and wounds in conjunction with skin sutures and stables or, after their removal, for wound support.
Reference	R150C, R1551C R150V, R1551V 801 5203.531
Basic UDI-DI	06082238401050000000044EA

are classified per rule(s) 4 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class1s 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: MDR 725202
Issued by: BSI, 2797

The Authorized European Representative for the concerned device(s) is

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

DocuSigned by:
Joe Coskey
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5/12/2026

Joe Coskey
Regulatory Manager
Consumer Business Group
3M Company

Date