EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Company
Single Registration Number *(TBD)*
2510 Conway Ave. St. Paul, MN 55144 USA

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

Trade Name*	Steri-Strip™ Skin Closures
Intended Purpose	Steri-Strip™ Skin Closures are indicated for use as a skin closure device in the treatment of lacerations and surgical incisions. Steri-Strip™ Adhesive Skin Closures may also be used in conjunction with skin sutures and staples or after their removal for wound support.
Reference	R1541NS, R1546NS, R1547NS, R1548NS
Basic UDI-DI	06082238401010000000054AH

are classified per rule 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.

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

3M Deutschland GmbH
Health Care Business
Single Registration Number (TBD)
Carl-Schurz-Str. 1
41453 Neuss, Germany

Dianne Gibbs, Division Regulatory Affairs Manager
3M Company
2510 Conway Ave. St. Paul, MN 55144 USA

7 July 2020
Date

3M and Steri-Strip is a trademark of 3M.

Issued to Authorized Representative (EC REP)

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