# *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 | Steri-Strip™ Blend Tone Skin Closures |
| Intended Purpose | Steri-Strip Skin Closures are intended for use as skin closure devices in the treatment of lacerations and surgical incisions. They may be used in conjunction with skin sutures and staples or after their removal for wound support. |
| Reference | B1551, B1557, B1559 |
| Basic UDI-DI | 06082238401010000000066AQ |

is classified per rule 4 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class Is 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

EU Representative Address
3M Deutschland GmbH
Health Care Business
Single Registration Number, DE-AR-000011642
Carl-Schurz-Str. 1
41453 Neuss, Germany

Dianne Gibbs, RAC
Regulatory Affairs Director
3M Medical Solutions Division

22 November 2021
Date

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

*Issued to Authorized Representative (EC REP)*