3M Health Care Business

3M Center
2510 Conway Ave, Bldg. 275-5W-06
St. Paul, MN 55144 U.S.A.
651 733 1110

Declaration of Conformity

As Legal Manufacturer
We, 3M Health Care Business,
2510 Conway Ave
St. Paul, MN 55144 USA

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

3M™ Steri-Strip™ Skin Closures (Reinforced)
R1540, R1541, R1542, R1546, R1547, R1548, R1549, R1540-01, R1540-02, R1540-12, 1540NP, 1541NP, R1541-01, R1541-02, R1541-12, R1542-01, R1542-12, R1546-01, R1547-12, R1546-12, 1546NP, R1547-01, 1547NP, 1551NP, 1541SP

3M™ Steri-Strip™ Blend Tone Skin Closures
B1550, B1551, B1553, B1557, B1559, B1551-02, B1551-12

3M™ Steri-Strip™ Elastic Skin Closures
E4540, E4541, E4541-12, E4542, E4546, E4547, E4548, E4549, 4541-12

3M™ Steri-Strip™ Wound Closure System
W8512, W8514, W8516

3M™ Steri-Strip™ Skin Closure Rack, 1555

3M™ Steri-Strip™ Compound Benzoin Tincture, C1544

3M™ Steri-Strip™ Skin Closures for Consumer
R150C (Nexcare), 5203.531 (M-Plast)

are classified, per rule 4 of Annex IX of the Medical Device Directive 93/42/EEC, as amended per 2007/47/EC, as Class Is, and

are in accordance with those aspects of Annex V relating to sterility in the manufacture of sterile devices and Annex VII and all other applicable provisions of the Directive 93/42/EEC, as amended per 2007/47/EC 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 CE 00493 delivered by BSI, 0086

EU Representative Address
3M Deutschland GmbH
Health Care Business
Carl-Schurz-Str. 1
41453 Neuss, Germany

Signature: Karen Little FOR MJW Date: 10 MARCH 2014

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3M Health Care
Vice President, Regulatory Affairs and Quality Assurance
3M Critical and Chronic Care Solutions Division

REG-DOC-05-000036, Steri-Strip Wound Closures, Version 7,
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