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 Company, 3M Health Care,
3M Center, 2510 Conway Ave, Bldg. 275-5W-06
St. Paul, MN 55144 USA

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

3M™ Tegaderm™ I.V. Transparent Film Dressing with Border
1610, 1650, 1655
3M™ Tegaderm™ Film Transparent Film Dressing with Border
1614, 1616
3M™ Tegaderm™ Film Transparent Film Dressing Frame Style
1622NP, 1622P, 1622IP, 1622SP, 1622W, 1622W/5, 1624NP, 1624P, 1624IP, 1624SP, 1624W, 1624W/5, 1624W/10, 1624P-10, 1626, 1626NP, 1626P, 1626IP, 1626SP, 1626W, 1626W/5, 1626W/10, 1626P-10, 1627, 1628, 1629, 1630, 1630NP, 1630P, 1630IP, 1630SP, 1630W/5, 1632P-10, 1634, 9505W, 9506W
3M™ Nexcare™ Tegaderm™ Transparent Dressing
T1012

is classified, per rule 4 of Annex IX of the Medical Device Directive 93/42/EEC, as amended per 2007/47/EC,
as a Class IIa device
and

is in accordance with Annex V and Annex VII of 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.

In addition, we declare that the above mentioned devices fulfil the applicable provisions of the Directive 93/42/EEC, as amended per 2007/47/EC.

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 Rittle Leigh Date: 15 JUNE 2016

Karen Rittle Leigh
3M Health Care
Division Regulatory Affairs Manager
Critical & Chronic Care Solutions Division

REG-DOC-05-129792, Tegaderm Transparent Film Dressings, Version 6,
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