



Declaration of Conformity

As Legal Manufacturer

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

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

3M™ Tegaderm™ Antimicrobial Transparent Dressing
and
3M™ Tegaderm™ Antimicrobial I.V. Advanced Securement Dressing
(containing chlorhexidine gluconate)
Product Numbers:
9124 and 9132

is classified,
per Rule 13 of Annex IX of the Medical Device Directive 93/42/EEC, as amended per 2007/47/EC
as a Class III device
and

is in accordance with Annex II 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 device fulfils the applicable provisions of the Directive 93/42/EEC,
as amended per 2007/47/EC.
and

are in conformity with the type described in the EC Type-Examination certificate reference number: CE 698003
and the full quality assurance certificate EC 02242
delivered by British Standards Institute (BSI), Notified Body Number 2797

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

Signature: _____

Dianne Gibbs
3M Health Care
Regulatory Affairs Director
3M Medical Solutions Division

Date: 24 March 2022

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REG-DOC-05-362965, Tegaderm Antimicrobial Transparent Dressing and Tegaderm Antimicrobial I.V. Advanced
Securement Dressing DOC, Version 2,
Status: Review, Release Date: <<Release Date>>

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