3M Health Care Business

3M Center
St. Paul, MN 55144-1000
651 733 1110

Declaration of Conformity

We, 3M Health Care,

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

3M™ Tegaderm™ + Pad Film Dressing with Nonadherent Pad
Product numbers:
3582, 3584, 3586, 3587, 3588, 3589, 3590, 3591, 3593
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 of Directive 93/42/EEC, as amended per 2007/47/EC
on the approximation of the laws of the 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 CE00493
delivered by BSI, 0086

This certificate is valid for devices originating from the following sites:

3M Brookings
601 22nd Ave. South
Brookings, South Dakota, 57006 USA

EU Representative Address
3M Medica
Zweigniederlassung der 3M Deutschland GmbH
Trading as "3M Health Care"
Hammfeldamm 11
D-41453 Neuss, Germany

Signature _____ Date: 11 Mar 2010

Kathryn W. Foran
3M Health Care
Regulatory Affairs and Quality Assurance
Skin & Wound Care Division

REG-DOC-05-132100, Tegaderm + Pad Dresing with Non-Adherent Pad, Version 1,
Status: Release, Release Date: 3/9/2010 1:51:52 PM