Declaration of Conformity

We
3M Health Care
hereby declare under our sole responsibility
that the CE marked products to which this declaration relates,

3M™ Tegaderm™ + Pad Transparent Dressing with Absorbent Pad
Or
3M™ Tegaderm™ + Pad Film Dressing with Nonadherent Pad

Product numbers:
3582NP, 3584NP, 3586NP, 3589NP, 3582P, 3584P, 3586P, 3589P, 3590P, 3591P

are classified,
according to the rule of Annex IX of the Medical Device Directive 93/42/EEC,
as Class IIa sterile devices
and

are in accordance with Annex V of Directive 93/42/EEC
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.

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

3M Poland Site Wroclaw
Kwidzynska 6
51-416 Wroclaw, Poland
EU Representative
3M Health Care, D41453, Neuss, Germany

Signature: Suzanne M. Danielson Date: September 29, 2008

Suzanne M. Danielson
Regulatory Affairs and Quality Director
Medical Division

REG-TF-US-05-000042, Tegaderm + Pad Tech File, Version 8.0,
Status: Released, Release Date: 29 Sep 2008 10:54