



Declaration of Conformity

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

Cavilon No Sting Barrier Film

Product numbers:

3346E- 28 ml bottle

3346P - 28 ml bottle,

are classified,

per rule 4 of Annex IX of the Medical Device Directive 93/42/EEC, as amended per 2007/47/EC,
as Class I devices

and

are in accordance with 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.

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

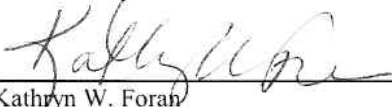
Zweigniederlassung der 3M Deutschland GmbH

Trading as "3M Health Care"

Hammfeldamm 11

D-41453 Neuss, Germany

Signature: _____


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

Date: _____

04-02-2010