



Declaration of Conformity

As Legal Manufacturer

We, 3M Company, 3M Health Care,
3M Center, 2510 Conway Ave,
Saint Paul, MN 55144 USA

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

3M™ Cavilon™ Advanced Skin Protectant
Product catalog numbers:
5050G

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, 2797

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

Signature: _____

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

Date: _____

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Page 1 of 1 Pages

Issued to St. Paul, Page 1 of 1 Pages