3M Health Care Business

3M Center, Building 02-1W-05-W-06
St. Paul, MN 55144-1000
651 733 1110

3M

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 Hydrogel Wound Filler (Unit Dose Hydrogel):
91110, 15 gram tube
91110-1, 15 gram tube
91111, 20 gram tube

is classified,
according to the rule of Annex IX of the Medical Device Directive 93/42/EEC,
as a Class II b medical device
and

is in accordance with Annex II of Directive 93/42/EEC 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.

This declaration is made on the basis of the quality assurance certificate CE 02242 delivered by BSI, 0086

This certificate is valid for the above devices placed on the market by
3M Healthcare
3M Center
Building 275
St. Paul, MN, USA 55144-1000

Signature: Suzanne M. Danielson Date: August 18, 2006

Suzanne M. Danielson
Regulatory Affairs and Quality Director
Medical Division