



***Declaration of Conformity***

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

**Micropore™ Surgical Tape**  
1530-0, 1530-1, 1530-2, 1530-3  
Single use  
1530S-1 and 1530S-2,  
Tan  
1533-0, 1533-1, 1533-2  
Dispenser  
1535-0, 1535-1, 1535-2, 1535-3

are classified, per Rule 1 of Annex IX of the Medical Device Directive 93/42/EEC, as amended per 2007/47/EC  
as a Class I device  
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.

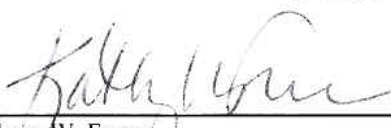
In addition, we declare that the above mentioned device fulfills the applicable provisions of the Directive  
93/42/EEC, as amended per 2007/47/EC.

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

3M Health Care  
3M Brookings  
601 22<sup>nd</sup> Ave. South,  
Brookings, South Dakota USA 57006

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: \_\_\_\_\_

12 Mar 2010