



Declaration of Conformity

As Legal Manufacturer
We, 3M Health Care,
2510 Conway Ave
St. Paul, MN 55144 U.S.A.

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

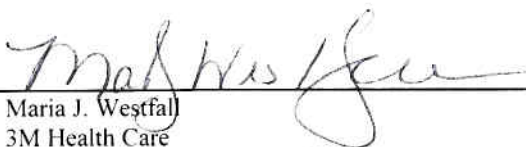
Micropore™ Surgical Tape (China)

1530-0 (CN), 1530-1 (CN), 1530-2 (CN),
1530-3 (CN), 1533-0 (CN), 1533-1 (CN), 1533-2 (CN),
1535-0 (CN), 1535-1(CN), 1535-2 (CN), 1535-3 (CN)

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.

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

Signature: 

Maria J. Westfal
3M Health Care
Vice President, Regulatory Affairs and Quality Assurance
Skin & Wound Care Division

Date: 6 August 2012