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## **Declaration of Conformity**

We

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

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

Micropore<sup>™</sup> Surgical Tape 1530-0, 1530-1, 1530-2, 1530-3 1530-0S, 1530-1S, 1530-2S, 1530-3S

Micropore<sup>™</sup> Plus+ Paper Tape 1532SS-1, 1532-2

are classified per rule 1 of Annex IX of the Medical Device Directive 93/42/EEC, as Class I devices

and

are in accordance with

Annex VII and all other applicable provisions of the Directive 93/42/EEC on the approximation of the laws of the European Union Member States concerning medical devices.

Signature:

Margaret Bessenbach

Manager Regulatory and Quality

Health Care Business West Europe & CEE 3M Deutschland GmbH, Health Care Business

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