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Document version: 3

Declaration of Conformity

We

**3M Poland Manufacturing Sp. z o.o.
ul. Kwidzyńska 6
51-416 Wrocław
Poland**

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

MICROPORE

(white)

1530-0, 1530-1, 1530-2, 1530-3
1530/1B, 1530/2B, 1530/5B, 1530/7B, 1530/10B
1530/1, 1530/2, 1530/5, 1530DR, 1530R, 1530-0/D, 1530-1/D,
1535-1, 1535E-0, 1535E-1, 1535E-2, 1535E-3, 1535E-4, 1535E-5
1530NP-0S, 1530NP-1S, 1530NP-0SD, 1530NP-1SD
1530SP-1, 1530SP-1D, 1530SP-0D, 1530N0, 1530N1, 1530N2

(tan)

1533/1B, 1533/2B, 1533/5B, 1533/2, 1533-P0/D, 1533-P1/D, 1533-1/D
1533NP-0SD, 1533NP-1SD, 1533NP-0S, 1533NP-1S
1533SP-1D, 1533SP-0D, 1533SP-1

Nexcare MICROPORE

NE-MIC125, NE-MIC250, NE-MIC500, NTM-1255, NTM-255, N1530-0D, N1530-1D

Nexcare SENSITIVE TAPE

N1530-1D

are classified, according to rules 1 of Annex IX of the
Medical Device Directive 93/42/EEC, as **Class I** non-sterile 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: _____

Marcin Baczyński
QMS/RA Supervisor

Date: 28.01.2016

relating to REG-STED-05-126457

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