



Document version: 16

Declaration of Conformity

We

**3M Poland Sp. Z o.o.
Al. Katowicka 117
Kajetany k/Warszawy
05-830 Nadarzyn
Poland**

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

MICROPORE

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, N1530-1D
1535E-0, 1535E-1, 1535E-2, 1535E-3, 1535E-4, 1535E-5
1530NP-0RCH, 1530NP-1RCH, 1530NP-0DCH, 1530NP-1DCH
NE-MIC125, NE-MIC250, NE-MIC500

MICROPORE (tan)

1533/1B, 1533/2B, 1533/5B, 1533/2, 1533-0/D, 1533-1/D
1533NP-0DCH, 1533NP-1DCH, 1533NP-0RCH, 1533NP-1RCH
1533-P1/D, 1533-P0/D

SENSITIVE TAPE

N1530-1D

are classified, according to rules 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: _____

Dominika Kawala
Quality Manager

Date: 24.02.2011

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