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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,

DURAPORE

**1538/5B, 1538-1/D, 1538NP-1SD
1538-0S, 1538-1S, 1538-2S**

POLOSILK

12,5 mm x 5 m; 25 mm x 5 m; 50 mm x 5 m

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: _____

02.02.2016

relating to REG-STED-05-207762

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