







(Full quality assurance system)

This is to certify that the company

## 3M Poland Manufacturing Sp. z.o.o.

ul. Kwidzynska 6 51-416 Wroclaw Poland

has implemented and maintains a full quality assurance system which applies to the products at every stage from design to final controls.

Through an audit, documented in a report, performed by DQS Medizinprodukte GmbH, it was verified that the management system fulfills the requirements of

## Annex II – excluding Section 4 of Council Directive 93/42/EEC concerning medical devices

with respect to the following medical devices:

Sterile Dressings class Is
Heating Plaster, class IIa
Hot and Cold Packs, class IIa
Blood Stop, class IIa
Sterile Gauze, class IIa
Sterile compresse gauze, class IIa
Sterile IV Cannula Dressing, class IIa

The manufacturer is subject to surveillance according to Annex II, Section 5. The CE marking with the Notified Body Identification Number (0297) may be affixed on the devices listed in the certificate. An EC Design Examination Certificate according to Annex II, Section 4 is required for class III devices covered by this certificate. The certificate is in the case of class I(s) devices (I(s) = class I products placed on the market in sterile conditions) limited to the aspects of manufacture concerned with securing and maintaining sterile conditions. The certificate is in the case of class I(m) devices (I(m) = class I devices with a measuring function) limited to the aspects of manufacture concerned with the conformity of the products with the metrological requirements.

Certificate registration no. 534977 MR2
Certificate unique ID 170712928
Effective date 2018-05-18
Expiry date 2023-03-16
Frankfurt am Main 2018-05-18

**DQS Medizinprodukte GmbH** 

Sigrid Uhlemann Managing Director Dr. Thomas Feldmann Head of Certification Body

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