

# EC CERTIFICATE

## for the Quality Assurance System



### according the Directive 93/42/EEC, Annex II excluding section (4)

As a Notified Body of the European Union, DEKRA Certification GmbH certifies, that the company  
**Murrplastik Medizintechnik GmbH**

Gewerbering 11, 08223 Falkenstein, Germany

**Certified location:**

Gewerbering 11, 08223 Falkenstein, Germany

applies a quality assurance system according to the Directive 93/42/EEC Annex II for the medical devices listed in the annex. The approval is based on the result of the re-certification audit report no. 51163-Z5-00, the decision dated 2018-12-18 and is only valid in connection with the successful performance of the annual surveillance audits.

This certificate is valid from 2018-12-20 to 2023-12-19

Registration No.: 51163-16-02



Ruth Delbeck-Bayer  
DEKRA Certification GmbH Stuttgart; 2018-12-18  
Notified Body ID-number: 0124

DEKRA Certification GmbH \* Handwerkstraße 15 \* D-70565 Stuttgart \* [www.dekra-certification.de](http://www.dekra-certification.de)



Benannt durch/Designated by  
Zentralstelle der Länder  
für Gesundheitsschutz  
bei Arzneimitteln und  
Medizinprodukten  
[www.zlg.de](http://www.zlg.de)  
**ZLG-BS-295.10.02**

# Annex to the EC Certificate No. 51163-16-02

Valid from 2018-12-20 to 2023-12-19

Revision status of the annex: 0 dated 2018-12-20

Devices/device categories included in the certificate:

## Class I s:

For the products listed below, review of the Quality Assurance System refers exclusively to aspects of manufacture concerned with securing and maintaining sterile conditions.

- MD 0101 + MDS 7006
  - Sterilset IIS

## Class II a:

- MD 0101 + MDS 7006
  - Y- Adapter 2.0// Y- Adapter 2.5// Y- Adapter 3.0// Y- Adapter 3.5
- MD 0101
  - Neofact Application aid



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