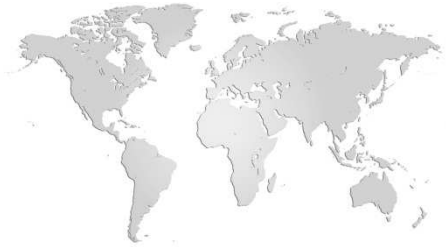
EC CERTIFICATE

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

j. söllner AG

Industriestraße 28, 94469 Deggendorf, 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. 51006-Z5-00, the decision dated 2021-05-17 and is only valid in connection with the successful performance of the annual surveillance audits.

This certificate is valid from 2021-05-17 to 2024-05-26

Registration No.: 51006-16-04

DEKRA Certification GmbH • DEKRA Certification • D-70565 Stuttgart, Handwerkstraße 15

Ruth Delbeck-Bayer
DEKRA Certification GmbH Stuttgart; 2021-05-17
Notified Body ID-number: 0124

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-295.10.02

Annex to the EC Certificate No. 51006-16-04

Valid from 2021-05-17 to 2024-05-26

Revision status of the annex: 0 dated 2021-05-17

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.

- Sterile tubing systems according rule 1
  - MD 0102 MDS 7006
- Sterile closures
  - MD 0102 MDS 7006
- Sterile adaptors
  - MD 0102 MDS 7006
- Sterile utilities
  - MD 0106 MDS 7006

Class II a:

- Sterile tubing systems according rule 2
  - MD 0102 MDS 7006

DEKRA Certification GmbH Stuttgart, Handwerkstraße 15 D-70565

Ruth Delbeck-Bayer
DEKRA Certification GmbH, Stuttgart, 2021-05-17
Notified Body ID-number: 0124