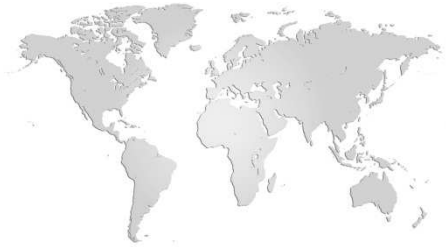
EC CERTIFICATE

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

MiE medical imaging electronics GmbH

Certified location:

Hauptstraße 112, 23845 Seth, 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. 50862-Z6-00, the decision dated 2019-07-15 and is only valid in connection with the successful performance of the annual surveillance audits.

This certificate is valid from 2019-07-16 to 2024-05-26

Registration No.: 50862-16-05

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

Ruth Delbeck-Bayer
DEKRA Certification GmbH Stuttgart; 2019-07-15
Notified Body ID-number: 0124

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
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