EC Certificate

Directive 93/42/EEC Annex II, excluding Section 4
Full Quality Assurance System
Medical Devices

Registration No.: HD 60147353 0001

Report No.: 21231982 015

Manufacturer:
Memmert GmbH + Co. KG
Äußere Rittersbacher Str. 38
91126 Schwabach
Deutschland

Products:
Dry Heat Sterilizers and CO2 Incubators
(see attachment for products and sites included)

Replaces Certificate, Registration No.: HD 60106200 0001

Expiry Date: 2024-05-26

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

Effective Date: 2020-02-27

Date: 2020-02-27

Notified Body

Roland Gruber

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.

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Doc. 1/1, Rev. 0

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Attachment to Certificate
Registration No.: HD 60147353 0001
Report No.: 21231982 015

Manufacturer:
Memmert GmbH + Co. KG
Äußere Rittersbacher Str. 38
91126 Schwabach
Deutschland

Products included:

Sterilizing Units Dry Heat, Type series S

CO2 Incubators
- ICO50med
- ICO105med
- ICO150med
- ICO240med

Site included:
- Memmert GmbH + Co. KG
  Willi-Memmert-Str. 90-96
  91186 Büchenbach, Germany

Activities: Production

Notified Body

Date: 2020-02-27

Roland Gruber

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle

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