

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

Registration No.: HD 60106200 0001

Report No.: 21231982 001

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 60099794 0001

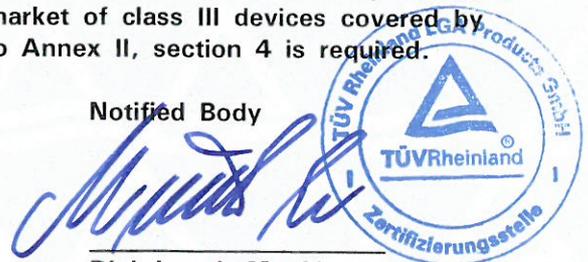
Expiry Date: 2020-12-07

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: 2015-12-08

Date: 2015-12-08

Notified Body



Dipl.-Ing. I. Munkler

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.

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

**Attachment to
Certificate**

Registration No.: HD 60106200 0001
Report No.: 21231982 005

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

Products included:

Sterilizing Units Dry Heat, Type series S

CO2 Incubators

- INCO108med
- INCO153med
- INCO246med
- ICO50med
- ICO105med
- ICO150med
- ICO240med

Site included:

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

Activities: Production

Date: 2017-08-16

Notified Body

Roland Grüber

